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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁷ : C12Q 1/70, 1/68, C12N 15/10, 15/34, 1/21, C07K 14/01, C12Q 1/18		A2	(11) International Publication Number: WO 00/32825 (43) International Publication Date: 8 June 2000 (08.06.00)																		
(21) International Application Number: PCT/IB99/02040 (22) International Filing Date: 3 December 1999 (03.12.99) (30) Priority Data: <table border="0"> <tr> <td>60/110,992</td> <td>3 December 1998 (03.12.98)</td> <td>US</td> </tr> <tr> <td>09/326,144</td> <td>3 June 1999 (03.06.99)</td> <td>US</td> </tr> <tr> <td>09/407,804</td> <td>28 September 1999 (28.09.99)</td> <td>US</td> </tr> <tr> <td>60/157,218</td> <td>30 September 1999 (30.09.99)</td> <td>US</td> </tr> <tr> <td>60/168,777</td> <td>1 December 1999 (01.12.99)</td> <td>US</td> </tr> <tr> <td>09/454,252</td> <td>2 December 1999 (02.12.99)</td> <td>US</td> </tr> </table> (71) Applicant (for all designated States except US): PHAGETECH, INC. [CA/CA]; Place du Parc, Case Postale 387, Montreal H2W 2N9 (CA). (72) Inventors; and (75) Inventors/Applicants (for US only): PELLETIER, Jerry [CA/CA]; 8 Lakeview, Baie D'Urfe, Quebec H9X 3B1 (CA). GROS, Phillippe [CA/CA]; 107 Montrose, St. Lambert, Quebec J4R 1X4 (CA). DUBOW, Michael [CA/CA]; 4901 Coolbrook Avenue, Montreal, Quebec H3X 2K8 (CA).		60/110,992	3 December 1998 (03.12.98)	US	09/326,144	3 June 1999 (03.06.99)	US	09/407,804	28 September 1999 (28.09.99)	US	60/157,218	30 September 1999 (30.09.99)	US	60/168,777	1 December 1999 (01.12.99)	US	09/454,252	2 December 1999 (02.12.99)	US	(74) Agents: MORROW, Joy, D. et al.; Smart & Biggar, 900 - 55 Metcalfe Street, P.O. Box 2999, Station D, Ottawa, Ontario K1P 5Y6 (CA). (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published Without international search report and to be republished upon receipt of that report.	
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(54) Title: DEVELOPMENT OF NOVEL ANTI-MICROBIAL AGENTS BASED ON BACTERIOPHAGE GENOMICS																					
(57) Abstract A method for identifying suitable targets for antibacterial agents based on identifying targets of bacteriophage-encoded proteins is described. Also described are compositions useful in the identification methods and in inhibiting bacterial growth, and methods for preparing and using such compositions.																					

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DESCRIPTION

Development of Novel Anti-Microbial Agents Based on Bacteriophage Genomics

5

BACKGROUND OF THE INVENTION

The present invention relates to the field of antibacterial agents and the treatment of infections of animals or other complex organisms by bacteria.

10 The frequency and spectrum of antibiotic-resistant infections have, in recent years, increased in both the hospital and community. Certain infections have become essentially untreatable and are growing to epidemic proportions in the developing world as well as in institutional settings in the developed world. The staggering spread of antibiotic resistance in pathogenic bacteria has been attributed to microbial
15 genetic characteristics, widespread use of antibiotic drugs, and changes in society that enhance the transmission of drug-resistant organisms. This spread of drug resistant microbes is leading to ever increasing morbidity, mortality and health-care costs.

 Ironically, it is the very success of antibiotics, resulting in their widespread use, that has contributed the most to rising numbers of drug resistant bacterial strains.
20 The longer a bacterial strain is exposed to a drug, the more likely it is to acquire resistance. Today, a total of 160 antibiotics, all based on a few basic chemical structures and targeting a small number of metabolic pathways, have found their way to market. Over-prescription of these drugs, as well as the failure of patients to comply with the complete antibiotic regimen, has lead to the rapid emergence of
25 antibiotic resistant strains. Such misuse of prescriptions, careless use of antibiotics in virtually all commercial production of beef and fowl, and changing societal conditions, such as the growth of day-care centers, increased long-term care in hospitals, and increased mobility of the population, has provided an environment where drug-resistant microbes can emerge and spread. Thus, virtually all common
30 infectious bacteria are becoming, or have already become, resistant to one or more groups of antibiotics. Such resistance now reaches all classes of antibiotics currently in use, including: β -lactams, fluoroquinolones, aminoglycosides, macrolide peptides, chloramphenicol, tetracyclines, rifampicin, folate inhibitors, glycopeptides, and
 mupirocin.

35 Over the last 45 years bacteria have adapted genetically to avoid the destruction/alteration of the essential pathways that these chemotherapeutic agents

target. Antibiotic resistant bacterial strains are now emerging at a higher rate than the rate at which new antibiotics are being developed. The consequence of this dilemma has been a dramatic increase in the cost of treating infections what would otherwise easily succumb to routine antibiotic therapy. Furthermore, and perhaps most
5 importantly, the emergence of multiple drug resistant pathogenic bacteria has led to a significant increase in morbidity and mortality, particularly in institutional settings.

Most major pharmaceutical companies have on-going drug discovery programs for novel anti-microbials. These are based on screens for small molecule inhibitors (natural products, bacterial culture media, libraries of small molecules,
10 combinatorial chemistry) of crucial metabolic pathways of the micro-organism of interest (*e.g.*, bacteria, fungi, parasites, worms). The screening process is largely for cytotoxic compounds and in most cases is not based on a known mechanism of action of the compounds. Pharmaceutical companies have large programs in this area. Classical drug screening programs are being exhausted and many of these
15 pharmaceutical companies are looking towards rational drug design programs.

Several small to mid-size biotechnology companies as well as large pharmaceutical companies have developed systematic high-throughput sequencing programs to decipher the genetic code of specific micro-organisms of interest. The goal is to identify, through sequencing, unique biochemical pathways or intermediates
20 that are unique to the microorganism. Knowledge of this may, in turn, form the rationale for a drug discovery program based on the mechanism of action of the identified enzymes/proteins. Genome Therapeutics Corp., The Institute for Genome Research, Human Genome Sciences Inc., and other companies have such sequencing programs in place. However, one of the most critical steps in this approach is the
25 ascertainment that the identified proteins and biochemical pathways are 1) non-redundant and essential for bacterial survival, and 2) constitute suitable and accessible targets for drug discovery.

SUMMARY OF THE INVENTION

While animals such as humans are, on occasion, infected by pathogenic bacteria, bacteria also have natural enemies. A number of host-specific viruses, known as bacteriophages or phages, infect and kill bacteria in the natural environment. Such bacteriophages generally have small compact genomes and bacteria are their exclusive hosts. Many known bacteria are host to a large number of bacteriophages that have been described in the literature. During the 1940's - 1960's, phage biology was an area of active research. As a testimony to this, the study of phages which infect and inhibit the enteric bacterium *Escherichia coli* (*E. coli*) contributed much to the early understanding of molecular biology and virology.

As is generally understood, bacteriophage (or phages) are viruses that infect and kill bacteria. They are natural enemies of bacteria and, over the course of evolution, have developed proteins (products of DNA sequences) which enable them to infect a host bacteria, replicate their genetic material, usurp host metabolism, and ultimately kill their host. The scientific literature well documents the fact that many known bacteria have a large number of such bacteriophages (Ackermann and DuBow, 1987) that can infect and kill them (for example, see the ATCC bacteriophage collection at <http://www.atcc.org>).

This invention utilizes the observation that bacteriophages successfully infect and inhibit or kill host bacteria, targeting a variety of normal host metabolic and physiological traits, some of which are shared by all bacteria, pathogenic and nonpathogenic alike. The term "pathogenic" as used herein denotes a contribution to or implication in disease or a morbid state of an infected organism. The invention thus involves identifying and elucidating the molecular mechanisms by which phages interfere with host bacterial metabolism, an objective being to provide novel targets for drug design. Whether the phage blocks bacterial RNA transcription or translation, or attacks other important metabolic pathways, such as cell wall assembly or membrane integrity, the basic blueprint for a phage's bacteria-inhibiting ability is encoded in its genome and can be unlocked using bioinformatics, functional genomics, and proteomics. By these means, the invention utilizes sequence information from the genomics of bacteriophage to identify novel antimicrobials that can be further used to actively and/or prophylactically treat bacterial infection.

Two important components of the invention thus are: i) the identification of bacteria-inhibiting phage open reading frames ("ORF"s) and corresponding products that can be used to develop antibiotics based on amino acid sequence and secondary structural characteristics of the ORF products, and ii) the use of bacteriophages to map

out essential bacterial target genes and homologs, which can in turn lead to the development of suitable anti-microbial agents. These two avenues represent new and general methods for developing novel antimicrobials.

The invention thus concerns the identification of bacteriophage ORFs that
5 supply bacteria-inhibiting functions. In this regard, use of the terms "inhibit",
"inhibition", "inhibitory", and "inhibitor" all refer to a function of reducing a
biological activity or function. Such reduction in activity or function can, for
example, be in connection with a cellular component, *e.g.*, an enzyme, or in
connection with a cellular process, *e.g.*, synthesis of a particular protein, or in
10 connection with an overall process of a cell, *e.g.*, cell growth. In reference to bacterial
cell growth, for example, an inhibitory effect (*i.e.*, a bacteria-inhibiting effect) may be
bacteriocidal (killing of bacterial cells) or bacteriostatic (*i.e.*, stopping or at least
slowing bacterial cell growth). The latter slows or prevents cell growth such that
fewer cells of the strain are produced relative to uninhibited cells over a given period
15 of time. From a molecular standpoint, such inhibition may equate with a reduction in
the level of, or elimination of, the transcription and/or translation of a specific
bacterial target(s), or reduction or elimination of activity of a particular target
biomolecule.

It is particularly advantageous to evaluate a plurality of different phage ORFs
20 for inhibitory activity that may be from one, but is preferably from a plurality of
different phage. For example, evaluating ORFs from a number of different phage of
the same bacterial host provides at least two advantages. One is that the multiple
phages will provide identification of a variety of different targets. Second, it is likely
that multiple phage will utilize the same cellular target

25 As used herein, the terms "bacteriophage" and "phage" are used
interchangeably to refer to a virus which can infect a bacterial strain or a number of
different bacterial strains.

In the context of this invention, the term "bacteriophage ORF" or "phage
ORF" or similar term refers to a nucleotide sequence in or from a bacteriophage. In
30 connection with a particular ORF, the terms refer an open reading frame which has at
least 95% sequence identity, preferably at least 97% sequence identity, more
preferably at least 98% sequence identity with an ORF from the particular phage
identified herein (*e.g.*, with an ORF as identified herein) or to a nucleic acid sequence
which has the specified sequence identify percentage with such an ORF sequence.

35 A first aspect of the invention thus provides a method for identifying a
bacteriophage nucleic acid coding region encoding a product active on an essential
bacterial target by identifying a nucleic acid sequence encoding a gene product which

provides a bacteria-inhibiting function when the bacteriophage infects a host bacterium, preferably one that is an animal or plant pathogen, more preferably a bird or mammalian pathogen, and most preferably a human pathogen. The bacteriophage is an uncharacterized bacteriophage. Thus, the method excludes, for example, phage λ , ϕ x174, m13 and other *E.coli*-specific bacteriophage that have been studied with respect to gene number and/or function. It also excludes, for example, the nucleic acid coding regions described in Tables 12-14, and in preferred embodiments, excludes the phage in which those regions are naturally located.

In connection with bacteriophage, the term "uncharacterized" means that a certain bacteriophage's genome has not yet been fully identified such that the genes having function involved in inhibiting host cells have not been identified. In particular, phage for which the description of genomic or protein sequence was first provided herein are uncharacterized. Phage sequences for which host bacteria-inhibiting functions have been identified prior to the filing of the present application (or alternatively prior to the present invention) are specifically excluded from the aspects involving utilization of sequences from uncharacterized bacteriophage, except that aspects may involve a plurality of phage where one or more of those phage are uncharacterized and one or more others have been characterized to some extent. A number of different bacteria-inhibiting phage ORFs are indicated in Tables 11-14. The phage ORFs or sequences identified therein are not within the term "uncharacterized; alternatively, in preferred embodiments the phage containing those ORFs are excluded from this term. Further, any additional phage ORFs (or alternatively the phage which contain those ORFs) which have previously been described in the art as bacteria-inhibiting ORFs are expressly excluded; those ORFs or phage are known to those skilled in the art and the exclusion can be made express by specifically naming such ORFs or phage as needed (likewise for uncharacterized targets as described below). For the sake of brevity, such a listing is not expressly presented, as such information is readily available to those skilled in the art.

Stating that an agent or compound is "active on" a particular cellular target, such as the product of a particular gene, means that the target is an important part of a cellular pathway which includes that target and that the agent acts on that pathway. Thus, in some cases the agent may act on a component upstream or downstream of the stated target, including on a regulator of that pathway or a component of that pathway.

By "essential", in connection with a gene or gene product, is meant that the host cannot survive without, or is significantly growth compromised, in the absence depletion, or alteration of functional product. An "essential gene" is thus one that encodes a product that is beneficial, or preferably necessary, for cellular growth in

vitro in a medium appropriate for growth of a strain having a wild-type allele corresponding to the particular gene in question. Therefore, if an essential gene is inactivated or inhibited, that cell will grow significantly more slowly, preferably less than 20%, more preferably less than 10%, most preferably less than 5% of the growth rate of the uninhibited wild-type, or not at all, in the growth medium. Preferably, in the absence of activity provided by a product of the gene, the cell will not grow at all or will be non-viable, at least under culture conditions similar to the *in vivo* conditions normally encountered by the bacterial cell during an infection. For example, absence of the biological activity of certain enzymes involved in bacterial cell wall synthesis can result in the lysis of cells under normal osmotic conditions, even though protoplasts can be maintained under controlled osmotic conditions. In the context of the invention, essential genes are generally the preferred targets of antimicrobial agents. Essential genes can encode target molecules directly or can encode a product involved in the production, modification, or maintenance of a target molecule.

15 A "target" refers to a biomolecule that can be acted on by an exogenous agent, thereby modulating, preferably inhibiting, growth or viability of a cell. In most cases such a target will be a nucleic acid sequence or molecule, or a polypeptide or protein. However, other types of biomolecules can also be targets, *e.g.*, membrane lipids and cell wall structural components.

20 The term "bacterium" refers to a single bacterial strain, and includes a single cell, and a plurality or population of cells of that strain unless clearly indicated to the contrary. In reference to bacteria or bacteriophage, the term "strain" refers to bacteria or phage having a particular genetic content. The genetic content includes genomic content as well as recombinant vectors. Thus, for example, two otherwise identical bacterial cells would represent different strains if each contained a vector, *e.g.*, a plasmid, with different phage ORF inserts.

In preferred embodiments, the phage is *Staphylococcus aureus* phage 77, 3A, 96, or 44 AHJD, *Enterococcus* sp. phage 182, or *Streptococcus pneumoniae* phage Dp-1.

30 In preferred embodiments, the phage is selected from. Preferred embodiments involve expressing at least one recombinant phage ORF(s) in a bacterial host followed by inhibition analysis of that host. Inhibition following expression of the phage ORF is indicative that the product of the ORF is active on an essential bacterial target. Such evaluation can be carried out in a variety of different formats, such as on a support matrix such as a solidified medium in a petri dish, or in liquid culture.

Preferably a plurality of phage ORFs are expressed in at least one bacterium. The plurality of phage ORFs can be from one or a plurality of phage. With respect to a single phage or at least one phage in a plurality of phages, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome. Preferably, for a plurality of phage, the plurality of expressed ORFs preferably represents at least 10%, more preferably at least 20%, 40%, or 60%, still more preferably at least 80% or 90%, and most preferably at least 95% of the ORFs in the phage genome of each phage. The plurality of phage ORFs can be expressed in a single bacterium, or in a plurality of bacteria where one ORF is expressed in each bacterium, or in a plurality of bacteria where a plurality of ORFs are expressed in at least one or in all of the plurality of bacteria, or combinations of these.

In embodiments of the above aspect (as well as in other aspects herein) in which a plurality of phage are utilized, a plurality of phage have the same bacterial host species; have different bacterial host species; or both. The plurality of phage includes at least two different phage, preferably at least 3,4,5,6,8,10,15,20, or more different phage. Indeed, more preferably, the plurality of phage will include 50, 75, 100, or more phage. As described herein, the larger number of phage is useful to provide additional target and target evaluation information useful in developing antibacterial agents, for example, by providing identification of a larger range of bacterial targets, and/or providing further indication of the suitability of a particular target (for example, utilization of a target by a number of different unrelated phage can suggest that the target is particularly stable and accessible and effective) and/or can indicate alternate sites on a target which interact with different inhibitors.

Further embodiments involve confirmation of the inhibitor function of the phage ORF, such as by utilizing or incorporating a control(s) designed to confirm the inhibitory nature of the ORF(s) being evaluated. The control can, for example, be provided by expression of an inactive or partially inactive form of the ORF or ORF product, and/or by the absence of expression of the ORF or ORF product in the same or a closely comparable bacterial strain as that used for expression of the test ORF. The reduced level of activity or the absence of active ORF product in the control will thus not provide the inhibition provided by a corresponding inhibitory ORF, or will provide a distinguishably lower level of inhibition. An inactivated or partially inactivated control has a mutation(s), *e.g.*, in the coding region or in flanking regulatory elements, that reduce(s) or eliminate(s) the normal function of the ORF. Thus, the inhibition of a bacterium following expression of a phage ORF is determined by comparison with the effects of expression of an inactivated ORF or the

response of the bacteria in the absence of expression in the same or similar type bacterium. Such determination of inhibition of the bacterium following expression of the ORF is indicative of a bacteria-inhibiting function. These manipulations are routinely understood and accomplished by those of skill in the art using standard techniques. In embodiments utilizing absence of expression of the ORF, the bacteria can, for example, contain an empty vector or a vector which allows expression of an unrelated sequence which is preferably non-inhibitory. Alternatively, the bacteria may have no vector at all. Combinations of such controls or other controls may also be utilized as recognized by those skilled in the art.

10 In embodiments involving expression of a phage ORF in a bacterial strain, in preferred embodiments that expression is inducible.

By "inducible" is meant that expression is absent or occurs at a low level until the occurrence of an appropriate environmental stimulus provides otherwise. For the present invention such induction is preferably controlled by an artificial environmental change, such as by contacting a bacterial strain population with an inducing compound (*i.e.*, an inducer). However, induction could also occur, for example, in response to build-up of a compound produced by the bacteria in the bacterial culture, *e.g.*, in the medium. As uncontrolled or constitutive expression of inhibitory ORFs can severely compromise bacteria to the point of eradication, such expression is therefore undesirable in many cases because it would prevent effective evaluation of the strain and inhibitor being studied. For example, such uncontrolled expression could prevent any growth of the strain following insertion of a recombinant ORF, thus preventing determination of effective transfection or transformation. A controlled or inducible expression is therefore advantageous and is generally provided through the provision of suitable regulatory elements, *e.g.*, promoter/operator sequences that can be conveniently transcriptionally linked to a coding sequence to be evaluated. In most cases, the vector will also contain sequences suitable for efficient replication of the vector in the same or different host cells and/or sequences allowing selection of cells containing the vector, *i.e.*, "selectable markers." Further, preferred vectors include convenient primer sequences flanking the cloning region from which PCR and/or sequencing may be performed.

As knowledge of the nucleotide sequence of phage ORFs is useful, *e.g.*, for assisting in the identification of phage proteins active against essential bacterial host targets, preferred embodiments involve the sequencing of at least a portion of the phage genome in combination with the above methods. This can be done either before or after or independent of expression and inhibition of the ORF in the bacteria, and provides information on the nature and characteristics of the ORF. Such a portion is

preferably at least 10%, 20%, 40%, 80%, 90%, or 100% of the phage genome. For embodiments in which a plurality of phage are utilized, preferably each phage is sequenced to an extent as just specified.

- Such sequencing is preferably accompanied by computer sequence analysis to
- 5 define and evaluate ORF(s), ORF products, structural motifs or functional properties of ORF products, and/or their genetic control elements. Thus, certain embodiments incorporate computer sequence analyses or nucleic acid and/or amino acid sequences. Further, existing data banks can provide phage sequence and product information which can be utilized for analysis and identification of ORFs in the sequence.
- 10 Computer analysis may further employ known homologous sequences from other species that suggest or indicate conserved underlying biochemical function(s) for the inhibitory or potentially inhibitory ORF sequence(s) being evaluated. This can include the sequences of signature motifs of identified classes of inhibitors.

- In the context of the phage nucleic acid sequences, e.g., gene sequences, of this
- 15 invention, the terms "homolog" and "homologous" denote nucleotide sequences from different bacteria or phage strains or species or from other types of organisms that have significantly related nucleotide sequences, and consequently significantly related encoded gene products, preferably having related function. Homologous gene sequences or coding sequences have at least 70% sequence identity (as defined by the
- 20 maximal base match in a computer-generated alignment of two or more nucleic acid sequences) over at least one sequence window of 48 nucleotides, more preferably at least 80 or 85%, still more preferably at least 90%, and most preferably at least 95%. The polypeptide products of homologous genes have at least 35% amino acid sequence identity over at least one sequence window of 18 amino acid residues, more
- 25 preferably at least 40%, still more preferably at least 50% or 60%, and most preferably at least 70%, 80%, or 90%. Preferably, the homologous gene product is also a functional homolog, meaning that the homolog will functionally complement one or more biological activities of the product being compared. For nucleotide or amino acid sequence comparisons where a homology is defined by a % sequence
- 30 identity, the percentage is determined using BLAST programs (with default parameters (Altschul et al., 1997, "Gapped BLAST and PSI-BLAST: a new generation of protein database search programs, *Nucleic Acid Res.* 25:3389-3402). Any of a variety of algorithms known in the art which provide comparable results can also be used, preferably using default parameters. Performance characteristics for
- 35 three different algorithms in homology searching is described in Salamov et al., 1999, "Combining sensitive database searches with multiple intermediates to detect distant

homologues." *Protein Eng.* 12:95-100. Another exemplary program package is the GCG™ package from the University of Wisconsin.

Homologs may also or in addition be characterized by the ability of two complementary nucleic acid strands to hybridize to each other under appropriately stringent conditions. Hybridizations are typically and preferably conducted with probe-length nucleic acid molecules, preferably 20-100 nucleotides in length. Those skilled in the art understand how to estimate and adjust the stringency of hybridization conditions such that sequences having at least a desired level of complementarity will stably hybridize, while those having lower complementarity will not. For examples of hybridization conditions and parameters, see, *e.g.*, Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. Homologs and homologous gene sequences may thus be identified using any nucleic acid sequence of interest, including the phage ORFs and bacterial target genes of the present invention.

A typical hybridization, for example, utilizes, besides the labeled probe of interest, a salt solution such as 6xSSC (NaCl and Sodium Citrate base) to stabilize nucleic acid strand interaction, a mild detergent such as 0.5% SDS, together with other typical additives such as Denhardt's solution and salmon sperm DNA. The solution is added to the immobilized sequence to be probed and incubated at suitable temperatures to preferably permit specific binding while minimizing nonspecific binding. The temperature of the incubations and ensuing washes is critical to the success and clarity of the hybridization. Stringent conditions employ relatively higher temperatures, lower salt concentrations, and/or more detergent than do non-stringent conditions. Hybridization temperatures also depend on the length, complementarity level, and nature (ie, "GC content") of the sequences to be tested. Typical stringent hybridizations and washes are conducted at temperatures of at least 40°C, while lower stringency hybridizations and washes are typically conducted at 37°C down to room temperature (~25°C). One of skill in the art is aware that these conditions may vary according to the parameters indicated above, and that certain additives such as formamide and dextran sulphate may also be added to affect the conditions.

By "stringent hybridization conditions" is meant hybridization conditions at least as stringent as the following: hybridization in 50% formamide, 5X SSC, 50 mM NaH₂PO₄, pH 6.8, 0.5% SDS, 0.1 mg/mL sonicated salmon sperm DNA, and 5X Denhart's solution at 42°C overnight; washing with 2X SSC, 0.1% SDS at 45°C; and washing with 0.2X SSC, 0.1% SDS at 45°C.

In sequence comparison analyses, an ORF, or motif, or set of motifs in a bacteriophage sequence can be compared to known inhibitor sequences, *e.g.*, homologous sequences encoding homologous inhibitors of bacterial function. Likewise, the analysis can include comparison with the structure of essential bacterial gene products, as structural similarities can be indicative of similar or replacement biological function. Such analysis can include the identification of a signature, or characteristic motif(s) of an inhibitor or inhibitor class.

Also, the identification of structural motifs in an encoded product, based on nucleotide or amino acid sequence analysis, can be used to infer a biochemical function for the product. A database containing identified structural motifs in a large number of sequences is available for identification of motifs in phage sequences. The database is PROSITE, which is available at www.expasy.ch/cgi-bin/scanprosite. The identification of motifs can, for example, include the identification of signature motifs for a class or classes of inhibitory proteins. Other such databases may also be used.

In aspects and preferred embodiments described herein, in which a bacterium or host bacterium is specified, the bacterium or host bacterium is preferably selected from a pathogenic bacterial species, for example, one selected from Table 1. Preferably, an animal or plant pathogen is used. For animals, preferably the bacterium is a bird or mammalian pathogen, still more preferably a human pathogen.

In aspects and preferred embodiments involving a bacteriophage or sequences from a bacteriophage, one or more bacteriophage are preferably selected from those listed in Table 1. Those exemplary bacteriophage are readily obtained from the indicated sources.

In some cases, it is advantageous to utilize phage with non-pathogenic host bacteria. The genome, structural motif, ORF, homolog, and other analyses described herein can be performed on such phage and bacteria. Such analysis provides useful information and compositions. The results of such analyses can also be utilized in aspects of the present invention to identify homologous ORFs, especially inhibitor ORFs in phage with pathogenic bacterial hosts. Similarly, identification of a target in a non-pathogenic host can be used to identify homologous sequences and targets in pathogenic bacteria, especially in genetically closely related bacteria. Those skilled in the art are familiar with bacterial genetic relationships and with how to determine relatedness based on levels of genomic identity or other measures of nucleotide sequence and/or amino acid sequence similarity, and/or other physical and culture characteristics such as morphology, nutritional requirements, or minimal media to support growth.

Also in preferred embodiments, an embodiment of this aspect is combined with an embodiment of the following aspect.

A related aspect of the invention provides methods for identifying a target for antibacterial agents by identifying the bacterial target(s) of at least one uncharacterized or untargeted inhibitor protein or RNA from a bacteriophage. Such identification allows the development of antibacterial agents active on such targets. Preferred embodiments for identifying such targets involve the identification of binding of target and phage ORF products to one another. The phage ORF products may be subportions of a larger ORF product that also binds the host target. In preferred embodiments, the phage protein or RNA is from an uncharacterized bacteriophage in Table 1. This aspect preferably includes the identification of a plurality of such targets in one or a plurality of different bacteria, preferably in one or a plurality of bacteria listed in Table 1.

In preferred embodiments of this aspect and other aspects of this invention involving particular phage ORFs or phage sequences, the ORF is *Staphylococcus aureus* phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As indicated for the above aspect, preferably the method involves the use of a plurality of different phage, and thus a plurality of different phage inhibitors and/or inhibitor ORFs.

In addition to uncharacterized phage ORF products, it is also useful to identify the targets of phage ORF products which are known to be inhibitors of host bacteria, but where the target has not been identified. Thus, such inhibitors can likewise be utilized as "untargeted" inhibitor phage ORFs and ORF products, e.g., proteins or RNAs.

In the context of inhibitor proteins or RNAs from a phage, the term "uncharacterized" means that a bacteria-inhibiting function for the protein has not previously been identified. Preferably, but not necessarily, the sequence of the protein or the corresponding coding region or ORF was not described in the art before the filing of the present application for patent (or alternatively prior to the present invention). Thus, this term specifically excludes any bacteria-inhibiting phage protein and its associated bacterial target which has been identified as inhibitory before the present invention or alternatively before the filing of the present application, for example those identified in Tables 12-14 or otherwise identified herein. For example, from *E. coli*, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, phage T4

gp55/gp33 alter the specificity of host RNA polymerase. The T4 *regB* gene product also targets the host translation apparatus. As with the uncharacterized bacteriophage ORFs or bacteriophage above, for such identified proteins, the sequences encoding those proteins are excluded from the uncharacterized inhibitor proteins.

5 The term "fragment" refers to a portion of a larger molecule or assembly. For proteins, the term "fragment" refers to a molecule which includes at least 5 contiguous amino acids from the reference polypeptide or protein, preferably at least 8, 10, 12, 15, 20, 30, 50 or more contiguous amino acids. In connection with oligo- or polynucleotides, the term "fragment" refers to a molecule which includes at least 15
10 contiguous nucleotides from a reference polynucleotide, preferably at least 24, 30, 36, 45, 60, 90, 150, or more contiguous nucleotides.

 Preferred embodiments involve identification of binding that include methods for distinguishing bound molecules, for example, affinity chromatography, immunoprecipitation, crosslinking, and/or genetic screen methods that permit
15 protein:protein interactions to be monitored. One of skill in the art is familiar with these techniques and common materials utilized (see, *e.g.*, Coligan, J. et al. (eds.) (1995) Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J.).

 Genetic screening for the identification of protein:protein interactions typically involves the co-introduction of both a chimeric bait nucleic acid sequence (here, the
20 phage ORF to be tested) and a chimeric target nucleic acid sequence that, when co-expressed and having affinity for one another in a host cell, stimulate reporter gene expression to indicate the relationship. A "positive" can thus suggest a potential inhibitory effect in bacteria. This is discussed in further detail in the Detailed Description section below. In this way, new bacterial targets can be identified that are
25 inhibited by specific phage ORF products or derivatives, fragments, mimetics, or other molecules.

 Other embodiments involve the identification and/or utilization of mutant targets by virtue of their host's relatively unresponsive nature in the presence of expression of ORFs previously identified as inhibitory to the non-mutant or wild-type
30 strain. Such mutants have the effect of protecting the host from an inhibition that would otherwise occur and indirectly allow identification of the precise responsible target for follow-up studies and anti-microbial development. In certain embodiments, rescue from inhibition occurs under conditions in which a bacterial target or mutant target is highly expressed. This is performed, for example, through coupling of the
35 sequence with regulatory element promoters, *e.g.*, as known in the art, which regulate expression at levels higher than wild-type, *e.g.*, at a level sufficiently higher that the

inhibitor can be competitively bound to the highly expressed target such that the bacterium is detectably less inhibited.

Identification of the bacterial target can involve identification of a phage-specific site of action. This can involve a newly identified target, or a target where the phage site of action differs from the site of action of a previously known antibacterial agent or inhibitor. For example, phage T7 genes 0.7 and 2.0 target the host RNA polymerase, which is also the cellular target for the antibacterial agent, rifampin. To the extent that a phage product is found to act at a different site than previously described inhibitors, aspects of the present invention can utilize those new, phage-specific sites for identification and use of new agents. The site of action can be identified by techniques well-known to those skilled in the art, for example, by mutational analysis, binding competition analysis, and/or other appropriate techniques.

Once a bacterial host target protein or nucleic acid or mutant target sequence has been identified and/or isolated, it too can be conveniently sequenced, sequence analyzed (e.g., by computer), and the underlying gene(s), and corresponding translated product(s) further characterized. Preferred embodiments include such analysis and identification. Preferably such a target has not previously been identified as an appropriate target for antibacterial action.

Certain embodiments include the identification of at least one inhibitory phage ORF or ORF product, e.g., as described for the above aspect, and thus are a combination of the two aspects.

Additionally, the invention provides methods for identifying targets for antibacterial agents by identifying homologs of a bacterial target e.g., *S. aureus*, *Enterococcus faecalis* or other *Enterococci*, and *Streptococcus pneumoniae* of a bacteriophage inhibitory ORF product. Such homologs may be utilized in the various aspects and embodiments described herein as described for the host *Enterococcus* sp. for bacteriophage 182.

Other aspects of the invention provide isolated, purified, or enriched specific phage nucleic acid and amino acid sequences, subsequences, and homologs thereof for phage selected from uncharacterized phage listed in Table 1, preferably from bacteriophage 77, 3A, 96, 44AHJD (*Staphylococcus aureus* host bacterium), Dp-1 (*Streptococcus pneumoniae* host), or 182 (*Enterococcus* host) or other phage listed in Table 1 for those bacteria. For example, such sequences do not include sequences identified in any of Tables 11-14. Nucleotide sequences of this aspect are at least 15 nucleotides in length, preferably at least 18, 21, 24, or 27 nucleotides in length, more preferably at least 30, 50, or 90 nucleotides in length. In certain embodiments, longer

nucleic acids are preferred, for example those of at least 120, 150, 200, 300, 600, 900 or more nucleotides. Such sequences can, for example, be amplification oligonucleotides (e.g., PCR primers), oligonucleotide probes, sequences encoding a portion or all of a phage-encoded protein, or a fragment or all of a phage-encoded protein. In preferred embodiments, the nucleic acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF. The upper length limit can also be expressed in terms of the number of base pairs of the ORF (coding region). In preferred embodiments, the nucleic acid sequence is from *Staphylococcus aureus* phage 77 ORF 17, 19, 43, 102, 104, or 182 as identified in U.S. application 09/407,804, *S. aureus* phage 44 AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As it is recognized that alternate codons will encode the same amino acid for most amino acids due to the degeneracy of the genetic code, the sequences of this aspect includes nucleic acid sequences utilizing such alternate codon usage for one or more codons of a coding sequence. For example, all four nucleic acid sequences GCT, GCC, GCA, and GCG encode the amino acid, alanine. Therefore, if for an amino acid there exists an average of three codons, a polypeptide of 100 amino acids in length will, on average, be encoded by 3^{100} , or 5×10^{47} , nucleic acid sequences. Thus, a nucleic acid sequence can be modified (e.g., a nucleic acid sequence from a phage as specified above) to form a second nucleic acid sequence encoding the same polypeptide as encoded by the first nucleic acid sequence using routine procedures and without undue experimentation. Thus, all possible nucleic acid sequences that encode the specified amino acid sequences are also fully described herein, as if all were written out in full, taking into account the codon usage, especially that preferred in the host bacterium. The alternate codon descriptions are available in common textbooks, for example, Stryer, BIOCHEMISTRY 3rd ed., and Lehninger, BIOCHEMISTRY 3rd ed., along with many others. Codon preference tables for various types of organisms are available in the literature. Sequences with alternate codons at one or more sites can also be utilized in the computer-related aspects and embodiments herein. Because of the number of sequence variations involving alternate codon usage, for the sake of brevity, individual sequences are not separately listed herein. Instead the alternate sequences are described by reference to the natural sequence with replacement of one or more (up to all e.g., up to 3, 5, 10, 15, 20, 30, 40, 50, or more) of the degenerate codons with alternate codons from the alternate codon

table (Table 6), or a modified table applicable to a particular organism that has differing codon usage, preferably with selection according to preferred codon usage for the normal host organism or a host organism in which a sequence is intended to be expressed. Those skilled in the art also understand how to alter the alternate codons to be used for expression in organisms where certain codons code differently than shown in the "universal" codon table.

For amino acid sequences or polypeptides, sequences contain at least 5 peptide-linked amino acid residues, and preferably at least 6, 7, 10, 15, 20, 30, or 40, amino acids having identical amino acid sequence as the same number of contiguous amino acid residues in a particular phage ORF product. In some cases longer sequences may be preferred, for example, those of at least 50, 60, 70, 80, or 100 amino acids in length. In preferred embodiments, the amino acid sequence contains a sequence which is within a length range with a lower length as specified above, and an upper length limit which is no more than 50, 60, 70, 80, or 90% of the length of the corresponding full-length ORF product. The upper length limit can also be expressed in terms of the number of amino acid residues of the ORF product. In preferred embodiments, the amino acid sequence or polypeptide has bacteria-inhibiting function when expressed or otherwise present in a bacterial cell which is a host for the bacteriophage from which the sequence was derived.

By "isolated" in reference to a nucleic acid is meant that a naturally occurring sequence has been removed from its normal cellular (*e.g.*, chromosomal) environment or is synthesized in a non-natural environment (*e.g.*, artificially synthesized). Thus, the sequence may be in a cell-free solution or placed in a different cellular environment. The term does not imply that the sequence is the only nucleotide chain present, but that it is essentially free (about 90-95% pure at least) of non-nucleotide material naturally associated with it, and thus is distinguished from isolated chromosomes.

The term "enriched" means that the specific DNA or RNA sequence constitutes a significantly higher fraction (2-5 fold) of the total DNA or RNA present in the cells or solution of interest than in normal or diseased cells or in cells from which the sequence was originally taken. This could be caused by a person by preferential reduction in the amount of other DNA or RNA present, or by a preferential increase in the amount of the specific DNA or RNA sequence, or by a combination of the two. However, it should be noted that enriched does not imply that there are no other DNA or RNA sequences present, just that the relative amount of the sequence of interest has been significantly increased.

The term "significant" is used to indicate that the level of increase is useful to the person making such an increase and an increase relative to other nucleic acids of about at least 2-fold, more preferably at least 5- to 10-fold or even more. The term also does not imply that there is no DNA or RNA from other sources. The other source DNA may, for example, comprise DNA from a yeast or bacterial genome, or a cloning vector such as pUC19. This term distinguishes from naturally occurring events, such as viral infection, or tumor type growths, in which the level of one mRNA may be naturally increased relative to other species of mRNA. That is, the term is meant to cover only those situations in which a person has intervened to elevate the proportion of the desired nucleic acid.

It is also advantageous for some purposes that a nucleotide sequence be in purified form. The term "purified" in reference to nucleic acid does not require absolute purity (such as a homogeneous preparation). Instead, it represents an indication that the sequence is relatively more pure than in the natural environment (compared to the natural level, this level should be at least 2-5 fold greater, *e.g.*, in terms of mg/mL). Individual clones isolated from a cDNA library may be purified to electrophoretic homogeneity. The claimed DNA molecules obtained from these clones could be obtained directly from total DNA or from total RNA. The cDNA clones are not naturally occurring, but rather are preferably obtained via manipulation of a partially purified naturally occurring substance (messenger RNA). The construction of a cDNA library from mRNA involves the creation of a synthetic substance (cDNA) and pure individual cDNA clones can be isolated from the synthetic library by clonal selection of the cells carrying the cDNA library. Thus, the process which includes the construction of a cDNA library from mRNA and isolation of distinct cDNA clones yields an approximately 10^6 -fold purification of the native message. Thus, purification of at least one order of magnitude, preferably two or three orders, and more preferably four or five orders of magnitude is expressly contemplated.

The terms "isolated", "enriched", and "purified" as respect nucleic acids, above, may similarly be used to denote the relative purity and abundance of polypeptides (multimers of amino acids joined one to another by α -carboxyl: α -amino group (peptide) bonds). These, too, may be stored in, grown in, screened in, and selected from libraries using biochemical techniques familiar in the art. Such polypeptides may be natural, synthetic or chimeric and may be extracted using any of a variety of methods, such as antibody immunoprecipitation, other "tagging" techniques, conventional chromatography and/or electrophoretic methods. Some of the above utilize the corresponding nucleic acid sequence.

As indicated above, aspects and embodiments of the invention are not limited to entire genes and proteins. The invention also provides and utilizes fragments and portions thereof, preferably those which are "active" in the inhibitory sense described above. Such peptides or oligopeptides and oligo or polynucleotides have preferred
5 lengths as specified above for nucleic acid and amino acid sequences from phage; corresponding recombinant constructs can be made to express the encoded same. Also included are homologous sequences and fragments thereof.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art.

10 In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Also, by having particular phage ORFs, e.g., the phage ORFs identified herein (e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described), other antimicrobial sequences from other bacteriophage sources can be identified and isolated using methods
15 described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage antimicrobial DNA segments from other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences that are
20 highly homologous. The bacteriophage segment from a specific phage, e.g., an antimicrobial DNA segment, can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on nucleic acid and/or amino acid sequence comparisons. As with identified inhibitory sequences, such homologous coding sequences and products can be used as
25 antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

The nucleotide and amino acid sequences identified herein are believed to be correct, however, certain sequences may contain a small percentage of errors, e.g., 1-5%. In the event that any of the sequences have errors, the corrected sequences can be
30 readily provided by one skilled in the art using routine methods. For example, the nucleotide sequences can be confirmed or corrected by obtaining and culturing the relevant phage, and purifying phage genomic nucleic acids. A region or regions of interest can be amplified, e.g., by PCR from the appropriate genomic template, using primers based on the described sequence. The amplified regions can then be
35 sequenced using any of the available methods (e.g., a dideoxy termination method).

This can be done redundantly to provide the corrected sequence or to confirm that the described sequence is correct. Alternatively, a particular sequence or sequences can be identified and isolated as an insert or inserts in a phage genomic library and isolated, amplified, and sequenced by standard methods. Confirmation or correction of a nucleotide sequence for a phage gene provides an amino acid sequence of the encoded product by merely reading off the amino acid sequence according to the normal codon relationships and/or expressed in a standard expression system and the polypeptide product sequenced by standard techniques. The sequences described herein thus provide unique identification of the corresponding genes, coding sequences, and other sequences, allowing those sequences to be used in the various aspects of the present invention.

In other aspects, the invention provides recombinant vectors and cells harboring at least one of the phage ORFs or portion thereof, or bacterial target sequences described herein. As understood by those skilled in the art, vectors may be provided in different forms, including, for example, plasmids, cosmids, and virus-based vectors. See, *e.g.*, Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; See also, Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J.

In preferred embodiments, the vectors will be expression vectors, preferably shuttle vectors that permit cloning, replication, and expression within bacteria. An "expression vector" is one having regulatory nucleotide sequences containing transcriptional and translational regulatory information that controls expression of the nucleotide sequence in a host cell. Preferably the vector is constructed to allow amplification from vector sequences flanking an insert locus. In certain embodiments, the expression vectors may additionally or alternatively support expression, and/or replication in animal, plant and/or yeast cells due to the presence of suitable regulatory sequences, *e.g.*, promoters, enhancers, 3' stabilizing sequences, primer sequences, etc. In preferred embodiments, the promoters are inducible and specific for the system in which expression is desired, *e.g.*, bacteria, animal, plant, or yeast. The vectors may optionally encode a "tag" sequence or sequences to facilitate protein purification. Convenient restriction enzyme cloning sites and suitable selective marker(s) are also optionally included. Such selective markers can be, for example, antibiotic resistance markers or markers which supply an essential nutritive growth factor to an otherwise deficient mutant host, *e.g.*, tryptophan, histidine, or leucine in the Yeast Two-Hybrid systems described below.

The term "recombinant vector" relates to a single- or double-stranded circular nucleic acid molecule that can be transfected into cells and replicated within or independently of a cell genome. A circular double-stranded nucleic acid molecule can be cut and thereby linearized upon treatment with appropriate restriction enzymes. An
5 assortment of nucleic acid vectors, restriction enzymes, and the knowledge of the nucleotide sequences cut by restriction enzymes are readily available to those skilled in the art. A nucleic acid molecule encoding a desired product can be inserted into a vector by cutting the vector with restriction enzymes and ligating the two pieces together. Preferably the vector is an expression vector, *e.g.*, a shuttle expression
10 vector as described above.

By "recombinant cell" is meant a cell possessing introduced or engineered nucleic acid sequences, *e.g.*, as described above. The sequence may be in the form of or part of a vector or may be integrated into the host cell genome. Preferably the cell is a bacterial cell.

15 In another aspect, the invention also provides methods for identifying and/or screening compounds "active on" at least one bacterial target of a bacteriophage inhibitor protein or RNA. Preferred embodiments involve contacting such a bacterial target or targets (*e.g.*, bacterial target proteins) with a test compound, and determining whether the compound binds to or reduces the level of activity of the bacterial target
20 (*e.g.*, a bacterial target protein). Preferably this is done either *in vivo* (*i.e.*, in a cell-based assay) or *in vitro*, *e.g.*, in a cell-free system under approximately physiological conditions.

The compounds that can be used may be large or small, synthetic or natural, organic or inorganic, proteinaceous or non-proteinaceous. In preferred embodiments,
25 the compound is a peptidomimetic, as described herein, a bacteriophage inhibitor protein or fragment or derivative thereof, preferably an "active portion", or a small molecule.

In preferred embodiments, the bacterial target is a target of a phage ORF identified herein, *e.g.*, *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus*
30 *pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

In particular embodiments, the methods include the identification of bacterial targets or the site of action of an inhibitor on a bacterial target as described above or otherwise described herein.

35 In embodiments involving binding assays, preferably binding is to a fragment or portion of a bacterial target protein, where the fragment includes less than 90%, 80%, 70%, 60%, 50%, 40%, or 30% of an intact bacterial target protein. Preferably,

the at least one bacterial target includes a plurality of different targets of bacteriophage inhibitor proteins, preferably a plurality of different targets. The plurality of targets can be in or from a plurality of different bacteria, but preferably is from a single bacterial species.

5 A "method of screening" refers to a method for evaluating a relevant activity or property of a large plurality of compounds (e.g., a bacteria-inhibiting activity), rather than just one or a few compounds. For example, a method of screening can be used to conveniently test at least 100, more preferably at least 1000, still more preferably at least 10,000, and most preferably at least 100,000 different compounds,
10 or even more.

In the context of this invention, the term "small molecule" refers to compounds having molecular mass of less than 2000 Daltons, preferably less than 1500, still more preferably less than 1000, and most preferably less than 600 Daltons. Preferably but not necessarily, a small molecule is not an oligopeptide.

15 In a related aspect or in preferred embodiments, the invention provides a method of screening for potential antibacterial agents by determining whether any of a plurality of compounds, preferably a plurality of small molecules, is active on at least one target of a bacteriophage inhibitor protein or RNA. Preferred embodiments include those described for the above aspect, including embodiments which involve
20 determining whether one or more test compounds bind to or reduce the level of activity of a bacterial target, and embodiments which utilize a plurality of different targets as described above.

The identification of bacteria-inhibiting phage ORFs and their encoded products also provides a method for identifying an active portion of such an encoded
25 product. This also provides a method for identifying a potential antibacterial agent by identifying such an active portion of a phage ORF or ORF product. In preferred embodiments, the identification of an active portion involves one or more of mutational analysis, deletion analysis, or analysis of fragments of such products. The method can also include determination of a 3-dimensional structure of an active
30 portion, such as by analysis of crystal diffraction patterns. In further embodiments, the method involves constructing or synthesizing a peptidomimetic compound, where the structure of the peptidomimetic compound corresponds to the structure of the active portion. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion that
35 the peptidomimetic will interact with the same molecule as the phage protein and preferably will elicit at least one cellular response in common which relates to the inhibition of the cell by the phage protein.

In preferred embodiments, the ORF or ORF product is or is derived or obtained from *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014 or product thereof.

5 The methods for identifying or screening for compounds or agents active on a bacterial target of a phage-encoded inhibitor can also involve identification of a phage-specific site of action on the target.

Preferably in the methods for identifying or screening for compounds active on such a bacterial target, the target is uncharacterized; the target is from an uncharacterized bacterium from Table 1; the site of action is a phage-specific site of action.

Further embodiments include the identification of inhibitor phage ORFs and bacterial targets as in aspects above.

15 An "active portion" as used herein denotes an epitope, a catalytic or regulatory domain, or a fragment of a bacteriophage inhibitor protein that is responsible for, or a significant factor in, bacterial target inhibition. The active portion preferably may be removed from its contiguous sequences and, in isolation, still effect inhibition.

By "mimetic" is meant a compound structurally and functionally related to a reference compound that can be natural, synthetic, or chimeric. In terms of the present invention, a "peptidomimetic," for example, is a compound that mimics the activity-related aspects of the 3-dimensional structure of a peptide or polypeptide in a non-peptide compound, for example mimics the structure of a peptide or active portion of a phage- or bacterial ORF-encoded polypeptide.

25 A related aspect provides a method for inhibiting a bacterial cell by contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein or RNA, where the target was uncharacterized. In preferred embodiments, the compound is such a protein, or a fragment or derivative thereof; a structural mimetic, *e.g.*, a peptidomimetic, of such a protein or fragment; a small molecule; the contacting is performed *in vitro*, the contacting is performed *in vivo* in an infected or at risk organism, *e.g.*, an animal such as a mammal or bird, for example, a human, or other mammal described herein; the bacterium is selected from a genus and/or species listed in Table 1; the bacteriophage inhibitor protein is uncharacterized; the bacteriophage inhibitor protein is from an uncharacterized phage listed in Table 1; the phage inhibitor protein is from one of *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

In the context of targets in this invention, the term "uncharacterized" means that the target was not recognized as an appropriate target for an antibacterial agent prior to the filing of the present application or alternatively prior to the present invention. Such lack of recognition can include, for example, situations where the target and/or a nucleotide sequence encoding the target were unknown, situations where the target was known, but where it had not been identified as an appropriate target or as an essential cellular component, and situations where the target was known as essential but had not been recognized as an appropriate target due to a belief that the target would be inaccessible or otherwise that contacting the cell with a compound active on the target *in vitro* would be ineffective in cellular inhibition, or ineffective in treatment of an infection. Methods described herein utilizing bacterial targets, *e.g.*, for inhibiting bacteria or treating bacterial infections, can also utilize "uncharacterized target sites", meaning that the target has been previously recognized as an appropriate target for an antibacterial agent, but where an agent or inhibitor of the invention is used which acts at a different site than that at which the previously utilized antibacterial agent, *i.e.*, a phage-specific site. Preferably the phage-specific site has different functional characteristics from the previously utilized site. In the context of targets or target sites, the term "phage-specific" indicates that the target or site is utilized by at least one bacteriophage as an inhibitory target and is different from previously identified targets or target sites.

In the context of this invention, the term "bacteriophage inhibitor protein" refers to a protein encoded by a bacteriophage nucleic acid sequence which inhibits bacterial function in a host bacterium. Thus, it is a bacteria-inhibiting phage product.

In the context of this invention, the phrase "contacting the bacterial cell with a compound active on a bacterial target of a bacteriophage inhibitor protein" or equivalent phrases refer to contacting with an isolated, purified, or enriched compound or a composition including such a compound, but specifically does not rely on contacting the bacterial cell with an intact phage which encodes the compound. Preferably no intact phage are involved in the contacting.

Related aspects provide methods for prophylactic or therapeutic treatment of a bacterial infection by administering to an infected, challenged or at risk organism a therapeutically or prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein or RNA, or as described for the previous aspect. Preferably the bacterium involved in the infection or risk of infection produces the identified target of the bacteriophage inhibitor protein or alternatively produces a homologous target compound. In preferred embodiments, the host organism is a plant or animal, preferably a mammal or bird, and more preferably, a human or other

mammal described herein. Preferred embodiments include, without limitation, those as described for the preceding aspect.

Compounds useful for the methods of inhibiting, methods of treating, and pharmaceutical compositions can include novel compounds, but can also include
5 compounds which had previously been identified for a purpose other than inhibition of bacteria. Such compounds can be utilized as described and can be included in pharmaceutical compositions.

In preferred embodiments of this and other aspects of the invention utilizing bacterial target sequences of a bacteriophage inhibitory ORF product, the target
10 sequence is encoded by a *Staphylococcus* nucleic acid coding sequence, preferably *S. aureus*, a *Streptococcus* nucleic acid coding sequence, preferably *Streptococcus pneumoniae*, or *Enterococcus* nucleic acid coding sequence. Possible target sequences are described herein by reference to sequence source sites.

The amino acid sequence of a polypeptide target is readily provided by
15 translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. For the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a
20 phage host genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

In the context of nucleic acid or amino acid sequences of this invention, the
25 term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the
30 homolog provides functionally equivalent biological function.

By "treatment" or "treating" is meant administering a compound or pharmaceutical composition for prophylactic and/or therapeutic purposes. The term "prophylactic treatment" refers to treating a patient or animal that is not yet infected but is susceptible to or otherwise at risk of a bacterial infection. The term "therapeutic
35 treatment" refers to administering treatment to a patient already suffering from infection.

The term "bacterial infection" refers to the invasion of the host organism, animal or plant, by pathogenic bacteria. This includes the excessive growth of bacteria which are normally present in or on the body of the organism, but more generally, a bacterial infection can be any situation in which the presence of a bacterial population(s) is damaging to a host organism. Thus, for example, an organism suffers from a bacterial population when excessive numbers of a bacterial population are present in or on the organism's body, or when the effects of the presence of a bacterial population(s) is damaging to the cells, tissue, or organs of the organism.

The terms "administer", "administering", and "administration" refer to a method of giving a dosage of a compound or composition, *e.g.*, an antibacterial pharmaceutical composition, to an organism. Where the organism is a mammal, the method is, *e.g.*, topical, oral, intravenous, transdermal, intraperitoneal, intramuscular, or intrathecal. The preferred method of administration can vary depending on various factors, *e.g.*, the components of the pharmaceutical composition, the site of the potential or actual bacterial infection, the bacterium involved, and the infection severity.

The term "mammal" has its usual biological meaning referring to any organism of the Class Mammalia of higher vertebrates that nourish their young with milk secreted by mammary glands, *e.g.*, mouse, rat, and, in particular, human, bovine, sheep, swine, dog, and cat.

In the context of treating a bacterial infection a "therapeutically effective amount" or "pharmaceutically effective amount" indicates an amount of an antibacterial agent, *e.g.*, as disclosed for this invention, which has a therapeutic effect. This generally refers to the inhibition, to some extent, of the normal cellular functioning of bacterial cells that renders or contributes to bacterial infection.

The dose of antibacterial agent that is useful as a treatment is a "therapeutically effective amount." Thus, as used herein, a therapeutically effective amount means an amount of an antibacterial agent that produces the desired therapeutic effect as judged by clinical trial results and/or animal models. This amount can be routinely determined by one skilled in the art and will vary depending on several factors, such as the particular bacterial strain involved and the particular antibacterial agent used.

In connection with claims to methods of inhibiting bacteria and therapeutic or prophylactic treatments, "a compound active on a target of a bacteriophage inhibitor protein" or terms of equivalent meaning differ from administration of or contact with an intact phage naturally encoding the full-length inhibitor compound. While an intact phage may conceivably be incorporated in the present methods, the method at

least includes the use of an active compound as specified different from a full length inhibitor protein naturally encoded by a bacteriophage and/or a delivery or contacting method different from administration of or contact with an intact phage encoding the full-length protein. Similarly, pharmaceutical compositions described herein at least
5 include an active compound different from a full-length inhibitor protein naturally encoded by a bacteriophage or such a full-length protein is provided in the composition in a form different from being encoded by an intact phage. Preferably the methods and compositions do not include an intact phage.

In accord with the above aspects, the invention also provides antibacterial
10 agents and compounds active on bacterial targets of bacteriophage inhibitor proteins or RNAs, where the target was uncharacterized as indicated above. As previously indicated, such active compounds include both novel compounds and compounds which had previously been identified for a purpose other than inhibition of bacteria. Such previously identified biologically active compounds can be used in
15 embodiments of the above methods of inhibiting and treating. In preferred embodiments, the targets, bacteriophage, and active compound are as described herein for methods of inhibiting and methods of treating. Preferably the agent or compound is formulated in a pharmaceutical composition which includes a pharmaceutically acceptable carrier, excipient, or diluent. In addition, the invention provides agents,
20 compounds, and pharmaceutical compositions where an active compound is active on an uncharacterized phage-specific site.

In preferred embodiments, the target is as described for embodiments of aspects above.

Likewise, the invention provides a method of making an antibacterial agent.
25 The method involves identifying a target of a bacteriophage inhibitor polypeptide or protein or RNA, screening a plurality of compounds to identify a compound active on the target, and synthesizing the compound in an amount sufficient to provide a therapeutic effect when administered to an organism infected by a bacterium naturally producing the target. In preferred embodiments, the identification of the target and
30 identification of active compounds include steps or methods and/or components as described above (or otherwise herein) for such identification. Likewise, the active compound can be as described above, including fragments and derivatives of phage inhibitor proteins, peptidomimetics, and small molecules. As recognized by those skilled in the art, peptides can be synthesized by expression systems and purified, or
35 can be synthesized artificially. In preferred embodiments the inhibitory phage ORF products is from *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus*

pneumoniae phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As indicated above, sequence analysis of nucleotide and/or amino acid sequences can beneficially utilize computer analysis. Thus, in additional aspects the invention provides computer-related hardware and media and methods utilizing and incorporating sequence data from uncharacterized phage, *e.g.*, uncharacterized phage listed in Table 1, preferably at least one of *Staphylococcus aureus* phage *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014, or 44 AHJD, *Enterococcus* sp. phage 182, or *Streptococcus pneumoniae* phage Dp-1. In general, such aspects can facilitate the above-described aspects. Various embodiments involve the analysis of genetic sequence and encoded products, as applied to the evaluating bacteriophage inhibitor ORFs and compounds and fragments related thereto. The various sequence analyses, as well as function analyses, can be used separately or in combination, as well as in preceding aspects and embodiments. Use in combination is often advantageous as the additional information allows more efficient prioritizing of phage ORFs for identification of those ORFs that provide bacteria-inhibiting function.

In one aspect, the invention provides a computer-readable device which includes at least one recorded amino acid or nucleotide sequence corresponding to one of the specified phage and a sequence analysis program for analyzing a nucleotide and/or amino acid sequence. The device is arranged such that the sequence information can be retrieved and analyzed using the analysis program. The analysis can identify, for example, homologous sequences or the indicated %s of the phage genome and structural motifs. Preferably the sequence includes at least 1 phage ORF or encoded product, more preferably at least 10%, 20%, 30%, 40%, 50%, 70%, 90%, or 100% of the genomic phage ORFs and/or equivalent cDNA, RNA, or amino acid sequences. Preferably the sequence or sequences in the device are recorded in a medium such as a floppy disk, a computer hard drive, an optical disk, computer random access memory (RAM), or magnetic tape. The program may also be recorded in such medium. The sequences can also include sequences from a plurality of different phage.

In this context, the term "corresponding" indicates that the sequence is at least 95% identical, preferably at least 97% identical, and more preferably at least 99% identical to a sequence from the specified phage genome, a ribonucleotide equivalent, a degenerate equivalent (utilizing one or more degenerate codons), or a homologous sequence, where the homolog provides functionally equivalent biological function.

Similarly, the invention provides a computer analysis system for identifying biologically important portions of a bacteriophage genome. The system includes a data storage medium, *e.g.*, as identified above, which has recorded thereon a nucleotide sequence corresponding to at least a portion of at least one uncharacterized bacteriophage genome, a set of program instructions to allow searching of the sequence or sequences to analyze the sequence, and an output device where the portion includes at least the sequence length as specified in the preceding aspect. The output device is preferably a printer, a video display, or a recording medium. More than one output device may be included. For each of the present computer-related aspects, the bacteriophage are preferably selected from the uncharacterized phage listed in Table 1, more preferably from bacteriophage 77, 3A, 96, 44 AHJD (*S. aureus*), Dp-1 (*Streptococcus pneumoniae*), or 182 (*Enterococcus*).

In keeping with the computer device aspects, the invention also provides a method for identifying or characterizing a bacteriophage ORF by providing a computer-based system for analyzing nucleotide or amino acid sequences, *e.g.*, as describe above. The system includes a data storage medium which has recorded a sequences or sequences as described for the above devices, a set of instructions as in the preceding aspect, and an output device as in the preceding aspect. The method further involves analyzing at least one sequence, and outputting the analysis results to at least one output device.

In preferred embodiments, the analysis identifies a sequence similarity or homology with a sequence or sequences selected from bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors; and essential bacterial ORFs. Preferably the analysis identifies a probable biological function based on identification of structural elements or characteristic or signature motifs of an encoded product or on sequence similarity or homology. Preferably the uncharacterized bacteriophage is from Table 1, more preferably at least one of bacteriophage 77, 3A, 96, 44 AHJD (*S. aureus*), Dp-1 (*Streptococcus pneumoniae*), or 182 (*Enterococcus*). In preferred embodiments, the method also involves determining at least a portion of the nucleotide sequence of at least one uncharacterized bacteriophage as indicated, and recording that sequence on data storage medium of the computer-based system. In preferred embodiments, the analysis identifies a sequence similarity of homology with a *S. aureus* phage 44AHJD ORF 1, 9, or 12, *Streptococcus pneumoniae* phage Dp-1 ORF 001, 002, 004, 008, 010, 013, 016, 021, 029, 030, 038, or 041, or *Enterococcus* sp. phage 182 ORF 002, 008, or 014.

As used in the claims to describe the various inventive aspects and embodiments, "comprising" means including, but not limited to, whatever follows the word "comprising". Thus, use of the term "comprising" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present. By "consisting of" is meant including, and limited to, whatever follows the phrase "consisting of". Thus, the phrase "consisting of" indicates that the listed elements are required or mandatory, and that no other elements may be present. By "consisting essentially of" is meant including any elements listed after the phrase, and limited to other elements that do not interfere with or contribute to the activity or action specified in the disclosure for the listed elements. Thus, the phrase "consisting essentially of" indicates that the listed elements are required or mandatory, but that other elements are optional and may or may not be present depending upon whether or not they affect the activity or action of the listed elements.

Further embodiments will be apparent from the following Detailed Description and from the claims.

BRIEF DESCRIPTION OF THE DRAWINGS

FIGURE 1A and 1B are flow schematics showing the manipulations used to convert pT0021, an arsenite inducible vector containing the luciferase gene, into pTHA or pTM, two *ars* inducible vectors. Vector pTHA contains BamHI, SalI, and HindIII cloning sites and a downstream HA epitope tag. Vector pTM contains BamHI and HindIII cloning sites and no HA epitope tag.

FIGURE 2 is a schematic representation of the cloning steps involved to place the DNA segments of any of ORFs 17/ 19/ 43/ 102/104/182 or other sequences into pTHA to assess inhibitory potential. For subcloning into pTM or pT0021, Individual ORFs were amplified by the PCR using oligonucleotides targeting the ATG and stop codons of the ORFs. Using this strategy, BamHI and HindIII sites were positioned immediately upstream or downstream, respectively of the start and stop codons of each ORF. Following digestion with BamHI and HindIII, the PCR fragments were subcloned into the same sites of pT0021 or pTM. Clones were verified by PCR and direct sequencing.

FIGURE 3 shows a schematic representation of the functional assays used to characterize the bactericidal and bacteriostatic potential of all predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Fig. 3A) Functional assay on semi-solid support media. Fig. 3B) Functional assay in liquid culture.

FIGURE 4A, B, and C is a bar graph showing the results of a screen in liquid media to assess bacteriostatic or bactericidal activity of 93 predicted ORFs (>33 amino acids) encoded by bacteriophage 77. Growth inhibition assays were performed as detailed in the Detailed Description. The relative growth of *Staphylococcus aureus* transformants harboring a given bacteriophage 77 ORF (identified on the bottom of the graph), in the absence or presence of arsenite, is plotted relative to growth of a *Staphylococcus aureus* transformant containing ORF 5, a non-toxic bacteriophage 77 ORF (which is set at 100%). Each bar represents the average obtained from three *Staph A* transformants grown in duplicate. Bacteriophage 77 ORFs showing significant growth inhibition consist of ORFs 17, 19, 102, 104, and 182.

FIGURE 5 shows a block diagram of major components of a general purpose computer.

FIGURE 6 shows an ORF map for *Streptococcus pneumoniae* bacteriophage Dp-1 showing the ORF identifiers, genomic locations, and orientations of the 85 identified ORFs that were found to have ribosomal binding sites and thus are expected to be expressed.

FIGURE 7 shows a schematic representation of the arsenite-inducible expression system present in a shuttle vector designed to express individual *Streptococcus* bacteriophage Dp-1 ORFs in *Streptococcus*. Various modifications can be readily made to such a vector, or other vectors can be readily constructed to provide inducible expression of ORFs in a particular host bacterium using well-known techniques.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention may be more clearly understood from the following description.

5 The tables will first be briefly described.

Table 1 is a listing of a large number of available bacteriophage that can be readily obtained and used in the present invention.

Table 2 shows the complete nucleotide sequence of the genome of *Staphylococcus aureus* bacteriophage 77.

10 Table 3 shows a list of all the ORFs from Bacteriophage 77 that were screened in the functional assay to identify those with anti-microbial activity.

Table 4 shows the predicted nucleotide sequence, predicted amino acid sequence, and physiochemical parameters of ORF 17/ 19/ 43/ 102/ 104/ 182]. These include the primary amino acid sequence of the predicted protein, the average
15 molecular weight, amino acid composition, theoretical pI, hydrophobicity map, and predicted secondary structure map.

Table 5 shows homology search results. BLAST analysis was performed with ORFs 17/ 19/ 43/ 102/ 104/ 182 against NCBI non-redundant nucleotide and Swissprot databases. The results of this search indicate that: I) ORF 17 has no
20 significant homology to any gene in the NCBI non-NCBI non-redundant nucleotide database, II) ORF 19 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 59 of bacteriophage phi PVL, III) ORF 43 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL, IV) ORF 102 has
25 significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 38 of phi PVL, V) ORF 104 has no significant homology to any gene in the NCBI non-redundant nucleotide database, VI) ORF 182 has significant homology to one gene in the NCBI non-redundant nucleotide database - the gene encoding ORF 39 of phi PVL.

30 Table 6 is a table from Alberts et al., MOLECULAR BIOLOGY OF THE CELL 3rd ed., showing the redundancy of the "universal" genetic code.

Table 7 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 3A.

Table 8 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 3A.

Table 9 shows the complete nucleotide sequence of *Staphylococcus aureus* bacteriophage 96.

5 Table 10 is a listing of the ORFs identified in *Staphylococcus aureus* bacteriophage 96.

Table 11 is a listing of sequences deposited in the NCBI public database (GeneBank) for bacteriophage listed in Table 1.

10 Table 12 is a listing of phage which encode a known lysis function , including the identified lysis gene.

Table 13 is a listing of bacteriophage which encode holin genes, where holin genes encode proteins which form pores and eventually enable other enzymes to kill the host bacterium.

Table 14 is a listing of bacteriophage which encode kil genes.

15 Table 15 is a list of *Staphylococcus aureus* sequences identified by accession number which may include sequences from genes coding for target sequences for the phage 77-encoded antimicrobial proteins or peptides. The sequences were obtained by searching GenBank for listings.

20 Table 16 shows the nucleotide sequence of the genome of *Staphylococcus aureus* phage 44 AHJD.

Table 17 lists and shows the sequence position of the 73 ORFs predicted to be encoded by *Staphylococcus aureus* bacteriophage 44 AHJD that are greater than 33 amino acids.

25 Table 18 shows the ORF sequences and putative amino acid sequences for the *Staphylococcus aureus* bacteriophage 44AHJD ORFs greater than 33 amino acids.

Table 19 shows the similarities in sequence identified between predicted *Staphylococcus aureus* bacteriophage 44 AHJD ORFs and sequences present in public databases.

30 Table 20 shows the homology alignments between predicted *Staphylococcus aureus* bacteriophage 44AHJD ORFs and the corresponding protein sequences present in public sequence databases.

Table 21 shows the complete nucleotide sequence of the genome of *Enterococcus* bacteriophage 182.

35 Table 22 lists and shows the sequence position of the 80 ORFs identified in bacteriophage 182 and that are greater than 33 amino acids.

Table 23 shows the nucleotide and predicted amino acid sequence of all 80 ORFs identified in bacteriophage 182.

Table 24 shows the similarities identified to date in sequence between *Enterococcus* phage 182 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 25 shows the predicted amino acid sequence as well as the predicted secondary structures map for two *Enterococcus* bacteriophage 182 ORFs.

Table 26 shows the homology alignments between predicted *Enterococcus* bacteriophage 182 ORFs and the corresponding protein sequences present in public sequence databases.

Table 27 list *Enterococcus* sequences listed in GenBank providing possible Enterococcal target sequences for inhibitory *Enterococcus* bacteriophage 182 ORFs and other compounds with antibacterial activity.

Table 28 shows the complete nucleotide sequence of the genome of *Streptococcus* bacteriophage Dp-1.

Table 29 lists and shows sequence position of the 273 ORFs identified in Pneumococcal bacteriophage Dp-1 that are greater than 33 amino acids, 85 of which are predicted to be expressed in Dp-1 as having a ribosomal binding site. That set of 85 ORFs is shown in the attached drawings.

Table 30 shows the nucleotide and predicted amino acid sequence of all 273 ORFs identified in bacteriophage Dp-1 that are identified as being expressed.

Table 31 shows the similarities identified in sequence between *Streptococcus* phage Dp-1 ORFs greater than 33 amino acids and sequences present in public sequence databases.

Table 32 shows the 4731 bp sequence of Dp-1 published by Sheehan et al., (1997).

Table 33 lists *Streptococcus pneumoniae* sequences listed in GenBank providing possible target sequences for inhibitory *Streptococcus pneumoniae* bacteriophage Dp-1 ORFs and other compounds with antibacterial activity

Background:

As indicated above, the present invention is concerned, in part, with the use of bacteriophage coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents. Thus, the invention concerns the selection of relevant bacteria. Particularly relevant bacteria are those which are pathogens of a complex organism such as an animal, e.g., mammals,

reptiles, and birds, and plants. Examples include *Staphylococcus aureus*, *Enterococcus* species, and *Streptococcus pneumoniae*. However, the invention can be applied to any bacterium (whether pathogenic or not) for which bacteriophage are available or which are found to have cellular components closely homologous to components targeted by phage of another bacterium.

Thus, the invention also concerns the bacteriophage which can infect a selected bacterium. Identification of ORFs or products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such targets are thus identified as potential targets for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, a phage-encoded inhibitor can also inhibit such a homologous bacterial cellular component.

The demonstration that bacteriophage have adapted to inhibiting a host bacterium by acting on a particular cellular component or target provides a strong indication that that component is an appropriate target for developing and using antibacterial agents, *e.g.*, in therapeutic treatments. Thus, the present invention provides additional guidance over mere identification of bacterial essential genes, as the present invention also provides an indication of accessibility of the target to an inhibitor, and an indication that the target is sufficiently stable over time (*e.g.*, not subject to high rates of mutation) as phage acting on that target were able to develop and persist. Thus, the present invention identifies a subset of essential cellular components which are particularly likely to be appropriate targets for development of antibacterial agents.

The invention also, therefore, concerns the development or identification of inhibitors of bacteria, in addition to the phage-encoded inhibitory proteins (or RNA transcripts), which are active on the targets of bacteriophage-encoded inhibitors. As described herein, such inhibitors can be of a variety of different types, but are preferably small molecules.

The following description provides preferred methods for use in the various aspects of the invention. However, as those skilled in the art will readily recognize, other approaches can be used to obtain and process relevant information. Thus the invention is not limited to the specifically described methods. In addition, the following description provides a set of steps in a particular order. That series of steps

describes the overall development involved in the present invention. However, it is clear that individual steps or portions of steps may be usefully practiced separately, and, further, that certain steps may be performed in a different order or even bypassed if appropriate information is already available or is provided by other sources or methods.

Selecting and Growing Phage, and Isolating DNA

Conceptually, the first step involves selecting bacterial hosts of interest. Preferably, but not necessarily, such hosts will be pathogens of clinical importance. Alternatively, because bacteria all share certain fundamental metabolic and structural features, these features can be targeted for study in one strain, for example a nonpathogenic one, and extrapolated to similarly succeed in pathogenic ones. Nonpathogenic strains may also exhibit initial advantages in being not only less dangerous, but also, for example, in having better growth and culturing characteristics and/or better developed molecular biology techniques and reagents. Consequently, advantageously the invention provides the ability target virtually any bacteria, but preferably pathogenic bacteria, with antimicrobial compounds designed and/or developed using bacteriophage inhibitory proteins and peptides from phage with non-pathogenic and/or pathogenic hosts.

We have selected *Staphylococcus aureus*, *Streptococcus pneumoniae*, various *Enterococci*, and *Pseudomonas aeruginosa* as initial exemplary pathogens. These bacteria are a major cause of morbidity and mortality in hospital-based infections, and the appearance of antibiotics resistance in all three organisms makes it increasingly difficult to treat benign infections involving these organisms. Such infections can include, for example, otitis media, sinusitis, and skin, and airway infections (Neu, H.C. (1992). *Science* 257, 1064-1073). However, the approach described below is clearly applicable to any human bacterial pathogens including but not restricted to *Mycobacterium tuberculosis*, *Nesseria gonorrhoeae*, *Haemophilus influenza*, *Acinobacter*, *Escherichia coli*, *Shigella dysenteria*, *Streptococcus pyogenes*, *Helicobacter pylori*, and *Mycoplasma* species. This invention can also be applied to the discovery of anti-bacterial compounds directed against pathogens of animals other than humans, for example, sheep, cattle, swine, dogs, cats, birds, and reptiles. Similarly, the invention is not limited to animals, but also applies to plants and plant pathogens.

In general, the bacteria are grown according to standard methodologies employed in the art, including solid, semi-solid or liquid culturing, which procedures can be found in or extrapolated from standard sources such as Maloy, S.R., Stewart,

V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press, or Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; or Ausubel, F.M. et al. (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. Culture conditions are selected which are adapted to the particular bacterium generally using culture conditions known in the art as appropriate, or adaptations of those conditions.

Nucleic acids within these bacteria can be routinely extracted through common procedures such as described in the above-referenced manuals and as generally known to those skilled in the art. Those nucleic acid stocks can then be used to practice the other inventive aspects described below.

Selection and Growth of Bacteriophage, and Isolation of DNA

The second step involves assembling a group of bacteriophages (phage collection) for one or more of the targeted bacterial hosts. While the invention can be utilized with a single bacteriophage for a pathogen or other bacterium, it is preferable to utilize a plurality of phage for each bacterium, as comparisons between a plurality of such phage provides useful additional information. Non-limiting examples of phage and sources for some of the above-mentioned pathogenic bacteria are found in Table 1. The criteria used to select such phages is that they are infectious for the microbe targeted, and replicate in, lyse, or otherwise inhibit growth of the bacterium in a measurable fashion. These phages can be very different from one another (representing different families), as judged by criteria such as morphology (head, tail, plate, etc.), and similarity of genome nucleotide sequence (cross-hybridization). Since such diverse bacteriophages are expected to block bacterial host metabolism and ultimately inhibit by a variety of mechanisms, their combined study will lead to the identification of different mechanisms by which the phages independently inhibit bacterial targets. Examples include degradation of host DNA (Parson K.A., and Snustad, D.P. (1975). *J. Virol.* 15, 221-444) and inhibition of host RNA transcription (Severinova, E., Severinov, K. and Darst, S.A. (1998). *J.Mol. Biol.* 279, 9-18). This, in turn, yields novel information on phage proteins that can inhibit the targeted microbe. As explained below, this 1) forms the basis of novel drug discovery efforts based on knowledge of the primary amino acid sequence of the phage inhibitor protein (e.g., peptide fragments or peptidomimetics) and/or 2) leads to the identification of bacterial biochemical pathways, the proteins of which are essential or significant for survival of the targeted microbe, and which enzymatic steps or

chemical reactions can be targeted by classical drug discovery methods using molecular inhibitors, for example, small molecule inhibitors.

Bacteriophage are generally either of two types, lytic or filamentous, meaning they either outright destroy their host and seek out new hosts after replication, or else continuously propagate and extrude progeny phage from the same host without destroying it. Regardless of the phage life cycle and type, preferred embodiments incorporate phage which impede cell growth in measurable fashion and preferably stop cell growth. To this end, lytic phage are preferred, although certain nonlytic species may also suffice, *e.g.*, if sufficiently bacteriostatic.

Various procedures that are commonly understood by those of skill in the art can be routinely employed to grow, isolate, and purify phage. Such procedures are exemplified by those found in such common laboratory aids such as Maloy, S.R., Stewart, V.J., and Taylor, R.K. Genetic Analysis of Pathogenic Bacteria (1996) Cold Spring Harbor Laboratory Press; Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.; and Ausubel, F.M. et al. (eds.) (1994) Current Protocols in Molecular Biology. John Wiley & Sons, Secaucus, N.J. The techniques generally involve the culturing of infected bacterial cells that are lysed naturally and/or chemically assisted, for example, by the use of an organic solvent such as chloroform that destroys the host cells thereby liberating the phage within. Following this, the cellular debris is centrifuged away from the supernatant containing the phage particles, and the phage then subsequently and selectively precipitated out of the supernatant using various methods usually employing the use of alcohols and/or other chemical compounds such as polyethylene glycol (PEG). The resulting phage can be further purified using various density gradient/centrifugation methodologies. The resulting phage are then chemically lysed, thereby releasing their nucleic acids that can be conveniently precipitated out of the supernatant to yield a viral nucleic acid supply of the phage of interest.

Exemplary bacteriophage are indicated in Table 1, along with sources where those phage may be obtained.

Exemplary bacteria include the reference bacteria for the identified bacteriophage, available from the same sources.

Characterizing Bacteriophage Genomes for ORFs

The third step involves systematically characterizing the genetic information contained in the phage genome. Within this genetic information is the sequence of all RNAs and proteins encoded by the phage, including those that are essential or

instrumental in inhibiting their host. This characterization is preferably done in a systematic fashion. For example, this can be done by first isolating high molecular weight genomic DNA from the phage using standard bacterial lysis methods, followed by phage purification using density gradient ultracentrifugation, and extraction of nucleic acid from the purified phage preparation. The high molecular weight DNA is then analyzed to determine its size and to evaluate a proper strategy for its sequencing. The DNA is broken down into smaller size fragments by sonication or partial digestion with frequently cutting restriction enzymes such as Sau3A to yield predominantly 1 to 2 kilobase length DNA, which DNA can then be resolved by gel electrophoresis followed by extraction from the gel.

The ends of the fragments are enzymatically treated to render them suitable for cloning and the pools of fragments are cloned in a bacterial plasmid to generate a library of the phage genome. Several hundred of these random DNA fragments contained in the plasmid vector are isolated as clones after introduction into an appropriate bacterium, usually *Escherichia coli*. They are then individually expanded in culture and the DNA from each individual clone is purified. The nucleotide sequences of the inserts of these clones are determined by standard automated or manual methods, using oligonucleotide primers located on either side of the cloning site to direct polymerase mediated sequencing (e.g., the Sanger sequencing method or a modification of that method). Other sequencing methods can also be used.

The sequence of individual clones is then deposited in a computer, and specific software programs (for example, Sequencher™, Gene Codes Corp.) are used to look for overlap between the various sequences, resulting in ordering of contig sequences and ultimately providing the complete sequence of the entire bacteriophage genome (one such example is given in Table 2 for *Staphylococcus aureus* bacteriophage 77; others are also provided herein). This complete nucleotide sequence is preferably determined with a redundancy of at least 3- to 5-fold (number of independent sequencing events covering the same region) in order to minimize sequencing errors.

Preferably, the bacterial strain used as a phage host should not possess any other innate plasmids, transposons, or other phage or incompatible sequences that would complicate or otherwise make the various manipulations and analyses more difficult.

Commercially available computer software programs are used to translate the nucleotide sequence of the phage to identify all protein sequences encoded by the phage (hereafter called open reading frames or ORFs). (Customized software can clearly also be used.) As phages are known to transcribe their genome into RNA from

both strands, in both directions, and sometimes in more than one frame for the same sequence, this exercise is done for both strands and in all six possible reading frames. As evolutionary constraints have forced the phage to conserve all of its vital protein sequences in as small a genome as possible, it is straightforward to identify all the proteins encoded by the phage by simple examination of the 6 translation frames of the genome. Once these ORFs are identified, they are cataloged into a phage proteome database (Table 3 lists ORFs identified from phage 77; ORF lists are also provided for other exemplary phage). This analysis is preferably performed for each phage under study. The process of ORF identification can be varied depending on the desired results. For example, the minimum length for the putative encoded polypeptide can be varied, and/or putative coding regions that have an associated Shine-Dalgarno sequence can be selected. In the case of phage 77 ORFs, such parameter adjustment was performed and resulted in the identification of ORFs as listed herein. Different parameters had resulted in the identification of the ORFs listed in the preceding U.S. Provisional Application 60/110,992, filed December 3, 1998, which is hereby incorporated by reference in its entirety.

Exemplary phage 77 ORFs identified in that provisional application and as identified herein are shown in the following table:

ORF ID from 60/110,992	Genomic position	a.a. size	Start codon	ORF ID from 241/190	Genomic position	a.a. size	Start codon
77ORF016	2369-24024	251	TTG	77ORF017	23269-23982	237	ATG
77ORF019	39845-40501	218	ATA	77ORF019	39851-40501	216	ATG
77ORF050	29268-29564	98	ATG	77ORF182	29268-29564	98	ATG
77ORF050	29268-29564	98	ATG	77ORF043	29304-29564	86	ATG
77ORF067	34312-34551	79	CTG	77ORF104	34393-34551	52	ATG
77ORF146	29051-29212	53	ATG	77ORF102	29051-29212	53	ATG

Identifying and Characterizing Inhibitory Phage ORFs

The fourth step entails identifying the phage protein or proteins or RNA transcripts that have the ability to inhibit their bacterial hosts. This can be accomplished, for example, by either or both of two non-mutually exclusive methods. The first method makes use of bioinformatics. Over the past few years, a large amount of nucleotide sequence information and corresponding translated products have become available through large genome sequencing projects for a variety of organisms including mammals, insects, plants, unicellular eukaryotes (yeast and fungi), as well as several bacterial genomes such as *E. coli*, *Mycobacterium tuberculosis*, *Bacillus subtilis*, *Staphylococcus aureus* and many others. Such sequences have been deposited in public databases (for example, non-redundant

sequence database at GenBank and SwissProt protein sequence database)
(<http://www.ncbi.nlm.nih.gov>) and can be freely accessed to compare any specific
query sequence to those present in such databases. For example, GenBank contains
over 1.6 billion nucleotides corresponding to 2.3 million sequence records. Several
5 computer programs and servers (e.g., TBLASTN) have been created to allow the rapid
identification of homology between any given sequence from one organism to that of
another present in such databases, and such programs are public and available free of
charge.

10 In addition, it has been well established that basic biochemical pathways can
be conserved in very distant organisms (for example bacteria and man), and that the
proteins performing the various enzymatic steps in these pathways are themselves
conserved at the amino acid sequence level. Thus, proteins performing similar
functions (e.g. DNA repair, RNA transcription, RNA translation) have frequently
preserved key structural signatures, identifiable by similarities across regions of
15 proteins (domains and motifs). The antimicrobials of the present invention will
preferably target features and targets that are highly characteristic or conserved in
microbes, and not higher organisms.

Most genomes encode individual proteins or groups of proteins that can be
assembled into protein families that have been evolutionarily conserved. Therefore,
20 similarity between a new query sequence and that of a member of a protein family
(reference sequences from public databases) can immediately suggest a biochemical
function for the novel query sequence, which in our case is a phage ORF.

The sequence homology between individual members of evolutionarily distant
members of a protein family is usually not randomly distributed along the entire
25 length of the sequence but is often clustered into "motifs" and "domains". These
correspond to key three-dimensional folds that form key catalytic and/or regulatory
structures that perform key biochemical function(s) for the group of proteins.
Commercially available computer software programs can identify such motifs in a
new query sequence, again providing functional information for the query sequence.
30 Such structural and functional motifs have also been derived from the combined
analysis of primary sequence databases (protein sequences) and protein structure
databases (X-ray crystallography, nuclear magnetic resonance) using so-called
"threading" methods (Rost B, and Sander C. (1996) *Ann. Rev. Biophys. Biomol.*
Struct. 25, 113-136).

35 Such motifs and folds are themselves deposited in public databases which can
be directly accessed (for example, SwissProt database; 3D-ALI at EMBL, Heidelberg;
PROSITE). This basic exercise leads to a structural homology map in which each of

the phage ORFs has been probed for such similarities, and where initial structural and functional hits are identified (selected examples of sequence homologies detected between individual ORFs from the genome of *Staphylococcus aureus* bacteriophage 77 and sequences deposited in public databases are shown in Table 5 for ORFs 17/19/43/102/104/182).

This analysis can point out phage proteins with similarity to proteins from other phages (such as those for *E. coli*) playing an important role in the basic biochemical pathways of the phage (such as DNA replication, RNA transcription, tRNAs, coat protein and assembly). Selected examples of such proteins include integrase and capsid protein. Therefore, this analysis enables identification and elimination of non-essential ORFs as candidates for an inhibitor function, as well as the identification of (potentially) useful ones.

In addition, this analysis can point out specific ORFs as possible inhibitor ORFs. For example these ORFs may encode proteins or enzymes that alter bacterial cell structure, metabolism or physiology, and ultimately viability. Examples of such proteins present in the genome of *Staphylococcus aureus* bacteriophage 77 include orf14 (deoxyuridine triphosphatase from bacteriophage T5), and orf15 (sialidase). (These ORF identifications are as listed in provisional application 60/110,992.) Other examples include ORFs 9 and 12 of *S. aureus* phage 44 AHJD, which encode the putative lysis functions found in many bacteriophages – a “holin” and an “amidase”.

In addition, it is well known that bacterial and eukaryotic viruses can usurp pathways from their host in order to use them to their advantage in blocking host cellular pathways upon infection. The phage can achieve this by 1) directly producing an inhibitor of a key host pathway (e.g. T7 gene 0.5 and 2), 2) directly producing a novel activity (e.g. T4 DNA polymerase), and 3) altering concentrations of cell components by producing similar functions (e.g. T4 transfer RNAs). The identification of sequence similarity between phage ORFs and bacterial host genome sequences will be highly indicative of such a mechanism. (Selected examples of such homologies are listed in Figure 4 of the provisional application 60/110,992 and include orf4 (homologous to autolysin), orf20 (hypothetical protein from *Staphylococcus aureus*) and orf29 (hypothetical protein from *Staphylococcus aureus*.) These ORFs can be analyzed by a standard biochemical approach to directly test their inhibitor functions (e.g., as described below).

Alternatively, a homology search may reveal that a given phage ORF is related to a protein present in the databases having an activity known to be inhibitory, (e.g., inhibitor of host RNA polymerase by *E. coli* bacteriophage T7. Such a finding would implicate the phage ORF product in a related activity. This will also suggest that a

new antimicrobial could be derived by a mimetic approach (*e.g.*, peptidomimetic) imitating this function or by a small molecule inhibitor to the bacterial target of the phage ORF, or any steps in the relevant host metabolic pathway, *e.g.*, high throughput screening of small molecule libraries. Selected examples of such similarity between
5 ORFs of *Staphylococcus aureus* bacteriophage 77 and proteins with inhibitor functions for bacterial hosts are listed in Figure 4 of the provisional application 60/110,992. These include orf9 (similar to bacteriophage P1 *kilA* function), and orf4 (autolysin of *Staphylococcus aureus*, amidase enzymatic activity).

A reason for the biochemical study of individual ORFs for inhibitor function is
10 that their expression or overexpression will block cellular pathways of the host, ultimately leading to arrest and/or inhibition of host metabolism. In addition, such ORFs can alter host metabolism in different ways, including modification of pathogenicity. Therefore, individual ORFs identified above are expressed, preferably overexpressed, in the host and the effect of this expression or overexpression on host
15 metabolism and viability is measured. This approach can be systematically applied to every ORF of the phage, if necessary, and does not rely on the absolute identification of candidate ORFs by bioinformatics. Individual ORFs are resynthesized from the phage genomic DNA, *e.g.*, by the polymerase chain reaction (PCR), preferably using oligonucleotide primers flanking the ORF on either side. These single ORFs are
20 preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial host such as *E. coli*, but containing the necessary information for plasmid replication in the target microbe such as *S. aureus* (hereafter referred to as shuttle vector). Shuttle vectors and their use are well known in the art.

Such shuttle vectors preferably also contain regulatory sequences that allow
25 inducible expression of the introduced ORF. As the candidate ORF may encode an inhibitor function that will eliminate the host, it is beneficial that it not be expressed prior to testing for activity. Thus, screening for such sequences when expressed in a constitutive fashion is less likely to be successful when the inhibitor is lethal. In the
30 exemplary inducible system presented in Figure 1A, 1B, 2, and 7, regulatory sequences from the *ars* operon of *S. aureus* are used to direct individual ORF expression in *S. aureus* (or other bacteria in which the *ars* system is functional). The *ars* operon encodes a series of proteins which normally mediate the extrusion of arsenite and other trivalent oxyanions from the cells when they are exposed to such
35 toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are

present. (Tauriainen, S. et al. (1997) *App. Env. Microb.*, Vol. 63, No. 11, p. 4456-4461.)

Therefore, individual phage ORFs can be expressed in *S. aureus* in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *S. aureus* clones expressing such individual phage ORFs. Toxicity of the phage inhibitor ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reduced or arrested host metabolism can be measured by pulse-chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis. Similar constructs can be made and used for other bacteria using well-known techniques.

Those skilled in the art are familiar with a variety of other inducible systems which can also be used for the controlled expression of phage ORFs, including, for example, lactose (see *e.g.*, Stratagene's LacSwitch™II system; La Jolla, CA) and tetracycline-based systems (see, *e.g.* Clontech's Tet On/Tet Off™ system; Palo Alto, CA). The arsenite-inducible system described is further depicted in Figures 1, 2 and 7.

The selection or construction of shuttle vectors and the selection and use of inducible systems are well known and thus other shuttle vectors appropriate for other bacteria can be readily provided by those skilled in the art, *e.g.*, for use in other bacterial species.

Standard methodologies for expressing proteins from constructs, and isolating and manipulating those proteins, for example in cross-linking and affinity chromatography studies, may be found in various commonly available and known laboratory manuals. See, *e.g.*, Current Protocols in Protein Science, John Wiley & Sons, Secaucus, N.J., and Maniatis, T. et al. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor University Press, Cold Spring, N.Y.

It has been found that certain phage or other viruses inhibit host cells, at least in part, by producing an antisense RNA which binds to and inhibits translation from a bacterial RNA sequence. Thus, in the case of potentially inhibitor RNA transcripts encoded by the phage genome, a strong indicator of a possible inhibitory function is provided by the identification of phage sequence which is identical to or fully complementary (or with only a small percentage of mismatch, *e.g.*, <10%, preferably less than 5%, most preferably less than 3%, to a bacterial sequence. This approach is convenient in the case of bacteria that have been essentially completely sequenced, as the comparison can be performed by computer using public database information.

The inhibitory effect of the transcript can be confirmed using expression of the phage sequence in a host bacterium. If needed, such inhibitory can also be tested by transfecting the cells with a vector that will transcribe the phage sequence to form RNA in such manner that the RNA produced will not be translated into a polypeptide.

5 Inhibition under such conditions provides a strong indication that the inhibition is due to the transcript rather than to an encoded polypeptide.

In an alternative, the expression of an ORF in a host bacterium is found to be inhibitory, but the inhibition is found to be due to an RNA product of the genomic coding region. For antisense inhibition, the sequence of the bacterial target nucleic acid sequence can be identified by inspection of the phage sequence, and the full sequence of the relevant coding region for the bacterial product can be found from a database of the bacterial genomic sequence or can be isolated by standard techniques (e.g., a clone in a genomic library can be isolated which contains the full bacterial ORF, and then sequenced).

10

15 In either case, the identification of a target which is inhibited by an RNA transcript produced by a phage provides both the possible inhibition of bacteria naturally containing the same target nucleic acid sequence, as well as the ability to use the target sequence in screening for other types of compounds which will act directly on the target nucleic acid sequence or on a polypeptide product expressed or regulated, at least in part, by the target of the inhibitory phage RNA.

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In some cases it will be found that the target of an inhibitory phage RNA or protein has previously been found to be a target of an inhibitory phage RNA or protein has previously been found to be a target for an antibacterial agent. In such cases, the phage inhibitor can still provide useful information if it is found that the phage-encoded product acts at a different site than the previously identified antibacterial agent or inhibitor, *i.e.*, acts at a phage-specific site. For many targets, action at a different site provides highly beneficial characteristics and/or information. For example, an alternate site of inhibitor action can at least partially overcome a resistance mechanism in a bacterium. As an illustration, in many cases, resistance is due, in large part, to altered binding characteristics of the immediate target to the antibacterial agent. The altered binding is due to a structural change which prevents or destabilizes the binding. However, the structural change is frequently quite local, so that compounds which bind at different local sites will be unaffected or affected to a much lesser degree. Indeed, in some cases the local sites will be on a different molecule and so may be completely unaffected by the local structural change creating resistance to the original agent(s). An example of resistance due to altered binding is

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provided by methicillin-resistant *Staphylococcus aureus*, in which the resistance is due to an altered penicillin-binding protein.

In other cases, a new site of action can have improved accessibility as compared to a site acted on by a previously identified agent. This can, for example, assist in allowing effective treatment at lower doses, or in allowing access by a larger range of types of compounds, potentially allowing identification of more potential active agents.

Another advantage is that the structural characteristics of a different site of action will lead to identification and/or development of inhibitors with different structures and different pharmacological parameter. This can allow a greater range of possibilities when selecting an antibacterial agent.

Yet further, different sites often produce different inhibitory characteristics in the target organism. This is commonly the case for multi-domain target proteins. Thus, inhibition targeting an alternate site can produce more efficacious action, e.g., faster killing, slower development of resistance, lower numbers of surviving cells, and different secondary effects (for example, different nutrient utilization).

Staphylococcus aureus phage 77

As indicated above, the present invention is concerned, in part, with the use of bacteriophage 77 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

As described, phage 77 ORFs 17, 19, 43, 102, 104, and 182 have been found to have bacteria inhibiting function. Identification of ORFs 17, 19, 43, 102, 104, and 182 and products from the phage which inhibit the host bacterium both provides an inhibitor compound and allows identification of the bacterial target affected by the phage-encoded inhibitor. Such a target is thus identified as a potential target for development of other antibacterial agents or inhibitors and the use of those targets to inhibit those bacteria. As indicated above, even if such a target is not initially identified in a particular bacterium, such a target can still be identified if a homologous target is identified in another bacterium. Usually, but not necessarily, such another bacterium would be a genetically closely related bacterium. Indeed, in some cases, an inhibitor encoded by phage 77 ORF 17, 19, 43, 102, 104, or 182 can also inhibit such a homologous bacterial cellular component.

Possible bacterial target sequences are described herein by reference to sequence source sites. In preferred embodiments, the sequence encoding the target corresponds

to a *S. aureus* nucleic acid sequence available from numerous sources including *S. aureus* sequences deposited in GenBank, *S. aureus* sequences found in European Patent Application No. 97100110.7 to Human Genome Sciences, Inc. filed January 7, 1997, *S. aureus* sequences available from TIGR at

- 5 <http://www.tigr.org/tdb/mdb/mdb.html>, and *S. aureus* sequences available from the Oklahoma University *S. aureus* sequencing project at the following URL: http://www.genome.ou.edu/staph_new.html. Such possible targets are particularly applicable to *S. aureus* phages 77, 3A, 96, and 44 AHJD.

- 10 The amino acid sequence of a polypeptide target is readily provided by translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a *S. aureus* coding sequence corresponding to a sequence listed in Table 15 herein. The listing in Table 15 describes *S. aureus* sequences currently listed with GenBank. Again, for the sake of brevity, the sequences are described by
- 15 reference to the database accession numbers instead of being written out in full herein. In cases where an entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage host *S. aureus* genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional
- 20 sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

Staphylococcus aureus phage 44 AHJD

- 25 The present invention also can utilize the identification of naturally occurring DNA sequence elements within *Staphylococcus aureus* bacteriophage 44AHJD which encode proteins with antimicrobial activity.

- Such identification can utilize bioinformatics identification of specific proteins (ORFs) utilized by *Staphylococcus aureus* bacteriophage 44AHJD during the viral life
- 30 cycle, resulting in a slowing or arrest of growth of the bacterial host, or in death, of the *Staphylococcus aureus* host including lysis of the infected bacteria. Thus, some of the bacteriophage 44AHJD DNA sequences encoding these proteins (ORFs) are predicted to encode antimicrobial functions. Information derived from these DNA sequences and translated ORFs can, in turn, be utilized to develop inhibitory
- 35 compounds by peptidomimetics that can also function as antimicrobials. In addition, the identification of the host bacterial proteins that are targeted and inhibited by the

antimicrobial bacteriophage ORFs can themselves provide novel targets for drug discovery.

The methodology described above is used to identify and characterize DNA sequences from *Staphylococcus* sp. bacteriophage 44 AHJD that have antimicrobial activity. As described in the Examples, the *Staphylococcus aureus* propagating strain (PS 44A), obtained from the Felix d'Herelle Reference Centre (#HER 1101), was used as a host to propagate its phage 44AHJD, also obtained from the Felix d'Herelle Reference Centre (#HER 101). By sequencing, we found that bacteriophage 44AHJD consists of 16,668 bp (Table 16) predicted to encode 73 ORFs greater than 33 amino acids (Tables 17 & 18). Computational analysis of the predicted protein products of *Staphylococcus aureus* bacteriophage 44AHJD identified homologs in public sequence databases as listed in Table 19 and 20, along with the accompanying list of related proteins.

From this analysis, it is apparent that 3 genes (ORF 3, 7, and 8) are related to structural proteins found in other bacteriophages. These include genes predicted to encode a tail protein (ORF 3), an upper collar/connector protein of the phage virion (ORF 7), and a lower collar protein (ORF 8). Bioinformatics has also identified one gene whose product is likely involved in phage DNA synthesis. One gene (ORF 1) shows significant homology to DNA polymerases of a number of bacteriophages, bacteria and fungi, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 44AHJD. ORF 2 encodes a protein with homology to the *dinC* gene of *Bacillus subtilis* that encodes a protein involved in teichoic acid biosynthesis. Teichoic acid is a polyphosphate polymer found in some, but not all, Gram positive organisms (and not in Gram negative organisms), where it is attached to the peptidoglycan layer. The phage protein may thus be involved in the synthesis of this material for incorporation into the cell wall, allowing enhanced lysis by the phage lysis enzymes or, as many enzymes can function in "reverse reactions", may be involved in its degradation allowing for penetration of the peptidoglycan and phage genome entry into the cell following adsorption. The similarity between *Staphylococcus aureus* bacteriophage 44AHJD and *E. coli* phage T7 indicates that they may share similar mechanisms of replication and growth. Both phages belong to the Podoviridae Family of bacteriophages and are members of the "T7-like" Genus of this Family (Ackermann and DuBow; VIth ICTV Report).

Two genes, ORF 9 and 12, were identified with the potential to encode antimicrobial protein products. The homology alignments are shown in Tables 19 and 20. The predicted product of ORF 9 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms, including that from the *Staphylococcus aureus* bacteriophage Twort. ORF 12 of *Staphylococcus aureus* bacteriophage 44AHJD shows homology to a set of lysis proteins from several bacteriophages. These lysis proteins are also referred to as holins, and represent phage-encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the cell wall and thus lyse the bacterium.

Thus, in particular embodiments, the present invention provides a nucleic acid sequence isolated from *Staphylococcus aureus* bacteriophage 44AHJD comprising at least a portion of one of the genes described above with antimicrobial activity. For example, ORF 1 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORF 9 directly encodes a polypeptide with antimicrobial activity. ORF 9 is predicted to encode an amidase, a protein known to act as a cell wall degrading enzyme. ORF 12 likely encodes a holin function required for transit of the phage amidase (gene 9 product) to the periplasm. When this type of gene product from Bacillus phage phi 29 (gene 14), was cloned in *Escherichia coli*, cell death ensued (Steiner et al., 1993). Thus, production of proteins from Bacillus phage phi 29 gene 14 in *E. coli* resulted in cell death, whereas production of protein from Bacillus phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner et al., 1993).

The present invention also provides the use of the *Staphylococcus* bacteriophage 44 AHJD antimicrobial ORFs or ORF products as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from *Staphylococcus* bacteriophage 44 AHJD killer ORFs.

Enterococcus phage 182

Bacteriophage 182 was obtained from the Felix D'Herelle phage collection (Ste. Foy, Quebec) and infects *Enterococcus* sp. Group D. The genome of
5 *Enterococcus* bacteriophage 182 consists of 17,833 bp (Table 21) and is predicted to encode 80 ORFs greater than 33 amino acids (Tables 22 and 23). Computational analysis of the predicted protein products of *Enterococcus* bacteriophage 182 was performed in order to identify protein products related to those deposited in public databases. Bacteriophage 182 protein products which detected sequences with
10 significant sequence similarity in public databases are listed in Table 24 and 26, along with the accompanying list of related proteins.

From this analysis, it is apparent that 5 genes (ORF 001, 004, 007, 009, and 011) are related to structural proteins of several *Bacillus* phages – *Bacillus* bacteriophage PZA, phi-29, and B103. These include genes predicted to encode a tail
15 protein (ORF 001), a head protein (ORF 004), and upper collar protein (ORF 007), a lower collar protein (ORF 009), and a pre-neck appendage protein (ORF 011). Two gene products are predicted to encode genes which direct phage morphogenesis – these are ORF 005 and 019.

Bioinformatics has also identified three genes whose products are likely
20 involved in phage DNA synthesis. One gene, ORF 002 shows significant homology to DNA polymerases of a number of bacteriophages, and the product of this gene is likely responsible for replicating the genetic material of bacteriophage 182. ORF 006 encodes a protein with homology to the encapsidation proteins of several other bacteriophages, including *Bacillus* phage phi-29 (P11014), PZA (P07541), and B103
25 (X99260) and *Streptococcus* phage CP-1 (Z47794). These gene products catalyze the *in vivo* and *in vitro* genome-encapsidation reaction (Garvey et al., 1985). Proteins involved in genome packaging have been shown to have additional activities that affect biochemical reactions in other phages and their hosts. For example, the coat protein of the RNA bacteriophage MS2 interacts with viral RNA to translationally
30 repress replicase synthesis (Pickett and Peabody, 1993). This protein-RNA interaction also plays a role in genome encapsidation, enveloping a single copy of the viral genome in a protein shell composed of many molecules of coat protein. In addition, the bacteriophage λ terminase enzyme can be lethal to *E. coli* when expressed,

suggesting cleavage of packaging sites in the bacterial chromosome. Also present within bacteriophage 182 is a gene, ORF 010, that encodes a protein that is related to the terminal proteins of *Bacillus* phage Nf (P06812), *Bacillus* phage GA-1 (X96987) and *Bacillus* phage B103 (X99260). DNA terminal proteins are linked to the 5' ends of both strands of the genome and are essential for DNA replication playing a role in initial priming of DNA replication. The similarity between *Enterococcus* bacteriophage 182 and *Bacillus* phages phi-29, PZA, and B103 indicates that they may share similar mechanisms of replication and growth. Protein-primed DNA replication is a well described phenomenon, and in the phi-29-like phages, the ends of the DNA serve as origins and termini of replication (Gutiérrez et al., 1986; Yoshikawa et al., 1985).

There is also a gene (ORF 015) that encodes a protein showing homology to an early protein product of *Bacillus* bacteriophage PZA and the single-strand nucleic acid binding protein of bacteriophage B103.

Two genes, ORF 008 and 014, were identified with the potential to encode anti-microbial protein products. The homology alignments are shown in Tables 24 & 26 and biochemical features of the predicted polypeptides shown in Table 25. The predicted product of ORF 008 is related to a class of genes which encodes lysozyme-like functions, enzymes which cleave linkages in the mucopolysaccharide cell wall structure of a variety of micro-organisms. ORF 014 of *Enterococcus* 182 shows homology to a set of lysis proteins from *Bacillus* bacteriophage phi-29, PZA, and B103. These lysis proteins are also referred to as holins and represent phage encoded lysis functions required for transit of the phage murein hydrolases (lysozyme) to the periplasm, where it can digest the outer cell wall and thus lyse the bacterium.

Thus, the present invention provides a nucleic acid sequence obtained from *Enterococcus* bacteriophage 182 comprising at least a portion of a phage 182 ORF, preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 002 encodes a DNA polymerase function. This polymerase may utilize host-derived accessory proteins for its activity when replicating the phage template, sequestering such proteins from use by the bacterial polymerase, resulting in inhibition of DNA replication, cell division, and cell growth. Alternatively, ORFs 008 or 014 directly encode polypeptides with anti-microbial activity. ORF 008 is predicted to encode an

autolytic lysozyme, a protein known to have anti-microbial activity (Martin *et al.*, 1998). ORF 014 likely encodes a holin function required for transit of the phage murein hydrolases to the periplasm. When the related product from *Bacillus* phage phi 29 (gene 14), was cloned in *Escherichia coli*, cell death ensued (Steiner *et al.*, 1993).
5 Thus, production of proteins from *Bacillus* phage phi 29 gene 14 in *E. coli* resulted in cell death, whereas production of protein from *Bacillus* phage phi 29 gene 14 concomitantly with the phi 29 lysozyme or unrelated murein-degrading enzymes led to lysis, suggesting that membrane-bound protein 14 induces a nonspecific lesion in the cytoplasmic membrane (Steiner *et al.*, 1993).

10 The present invention also provides the use of the *Enterococcus* bacteriophage 182 anti-microbial ORFs as pharmacological agents, either wholly or in part and derivatives, as well as the use of corresponding peptidomimetics, developed from amino acid or nucleotide sequence knowledge derived from *Enterococcus* bacteriophage 182 killer ORFs. This can be done where the structure of the
15 peptidomimetic compound corresponds to the structure of the active portion of a product of an ORF. In this analysis, the peptide backbone is transformed into a carbon based hydrophobic structure that can retain cytostatic or cytotoxic activity for the bacterium. This is done by standard medicinal chemistry methods, measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These
20 mimetics also represent lead compounds for the development of novel antibiotics. In this context, "corresponds" means that the peptidomimetic compound structure has sufficient similarities to the structure of the active portion of a product of one of the *Enterococcus* ORFs listed, that the peptidomimetic will interact with the same molecule as the product of the ORF, and preferably will elicit at least one cellular
25 response in common which relates to the inhibition of the cell by the phage protein.

To validate the identity of an ORF as a killer ORF, it is preferably expressed in the host or other test bacterial organism and the effect of this expression on bacterial growth and replication is assessed. Therefore, all individual ORFs identified herein, e.g., those identified above, can be expressed, preferably overexpressed, in a
30 suitable host bacterium e.g., a host *Enterococcus* and the effect of this expression or overexpression on host metabolism and viability can be measured.

Individual ORFs can be resynthesized from the phage genomic DNA by the polymerase chain reaction (PCR) using oligonucleotide primers flanking the ORF on

either side. Those skilled in the art are familiar with the design and synthesis of appropriate primer sequences. These single ORFs are preferably engineered so that they contain appropriate cloning sites at their extremities to allow their introduction into a new bacterial expression plasmid, allowing propagation in a standard bacterial
5 host such as *E. coli*, but containing the necessary information for plasmid replication in the target microbe, *Enterococcus* sp. (hereafter referred to as a shuttle vector).

This shuttle vector also preferably contains regulatory sequences that allow inducible expression of the introduced ORF. As the candidate ORF may encode a killer function that will eliminate the host, it is highly advantageous that it not be
10 expressed (or at least not expressed at a substantial level) prior to testing for activity; thus screening for such sequences in a constitutive fashion is less likely to be successful (lethality). In an example presented in Fig. 7, regulatory sequences from the *ars* operon are used to direct individual ORF expression in *Enterococcus*. The *ars* operon encodes a series of proteins which normally mediate the extrusion of arsenite
15 and several other trivalent oxyanions from the cells when they are exposed to such toxic substances in their environment. The operon encoding this detoxifying mechanism is normally silent and only induced when arsenite-related compounds are present.

Therefore, individual phage ORFs can be expressed in *Enterococcus* or other
20 suitable host in an inducible fashion by adding to the culture medium non-toxic arsenite concentrations during the growth of individual *Enterococcus* (or other host cells) clones expressing such individual phage ORFs. Toxicity of the phage killer ORF for the host is monitored by reduction or arrest of growth under induction conditions, as measured by optical density in liquid culture or after plating the
25 induced cultures on solid medium. Subsequently, interference of the phage ORF with the host biochemical pathways ultimately leading to reducing or arresting host metabolism can be measured by pulse chase experiments using radiolabeled precursors of either DNA replication, RNA transcription, or protein synthesis.

Of course, other inducible regulatory sequences (e.g., promoters, operators,
30 etc.) may be used (e.g., systems using positive induction of expression or systems using release of repression). A variety of such systems are known to those skilled in the art and can be utilized in the present invention.

Nucleic acid sequences of the present invention can be isolated using a method similar to those described herein or other methods known to those skilled in the art. In addition, such nucleic acid sequences can be chemically synthesized by well-known methods. Having the phage 182 ORFs, e.g., anti-bacterial ORFs of the present invention, portions thereof, or oligonucleotides derived therefrom as described, other
5 anti-microbial sequences from other bacteriophage sources can be identified and isolated using methods described here or other methods, including methods utilizing nucleic acid hybridization and/or computer-based sequence alignment methods.

The invention also provides bacteriophage anti-microbial DNA segments from
10 other phages based on nucleic acids and sequences hybridizing to the presently identified inhibitory ORF under high stringency conditions or sequences which are highly homologous. The bacteriophage anti-microbial DNA segment from bacteriophage 182 can be used to identify a related segment from another unrelated phage based on stringent conditions of hybridization or on being a homolog based on
15 nucleic acid and/or amino acid sequence comparisons. As with the phage 182 inhibitory sequences, such homologous coding sequences and products can be used as antimicrobials, to construct active portions or derivatives, to construct peptidomimetics, and to identify bacterial targets.

Enterococcus sequences are listed in Table 27 by accession number, providing
20 identification of possible targets of *Enterococcus* phage inhibitory ORF products, e.g., from phage 182.

Streptococcus pneumoniae

As indicated in the Summary above, the present invention is concerned
25 with the use of *Streptococcus* sp. bacteriophage Dp-1 coding sequences and the encoded polypeptides or RNA transcripts to identify bacterial targets for potential new antibacterial agents.

Streptococcus pneumoniae is an important cause of community-acquired pneumonia and a major cause of otitis media, sinusitis, and meningitis in children and
30 adults. In Spain and other Mediterranean countries, the majority of *S. pneumoniae* are relatively resistant to penicillin (Klugman, 1990; Fenoli et al., 1991; Jørgensen et al., 1990). These strains also have decreased susceptibility to broad-spectrum cephalosporins, which are frequently used in the empiric treatment of meningitis and

other serious invasive bacterial infections. High-level resistance of pneumococci has been encountered in Hungary where 70% of children who were colonized with *S. pneumoniae* carried penicillin resistant strains that were also resistant to tetracycline, erythromycin, trimethoprim/sulfamethoxazole, and 30% resistant to chloramphenicol (Neu, 1992). The resistance of pneumococci to macrolides such as erythromycin averages 20-25% in France, ~20% in Japan, and <10% in Spain (Neu, 1992).

The antimicrobial susceptibilities and distribution of serotypes of the 42 isolates of *S. pneumoniae* in southern Taiwan from invasive infections have been recently determined (Hseuh et al., 1996). Resistance rates among these isolates were: erythromycin, 61.9%; clindamycin, 47.6%; chloramphenicol, 19%; and tetracycline, 73.8%. Resistance to three or more classes of antibiotics was found in 33.3% of the isolates. Bacteremic pneumonia and primary bacteremia accounted for 64.3% of the infections and mortality was 42.6%. Given the severity of these infections despite adequate antibiotic therapy, there is clearly a need for introduction of new therapeutic options to prevent mortality due to invasive *S. pneumoniae* infections.

Pneumococcal phages belong to four families and they present a great variety in morphology, including lytic and temperate phages (for a review, see Garcia et al., 1997). Examples of lytic phages are Cp-1 and Dp-1, whereas examples of temperate phages are HB-3, EJ-1, and HB-746. The complete nucleotide sequence and functional organization of Cp-1 has been reported (Martin et al., 1996). Cp-1 has a 19,345 bp double-stranded DNA genome, with a terminal protein covalently linked to its 5' ends, that replicates by a protein primed mechanism. The phage contains 29 ORFs, 23 on one strand and 6 on the opposite. When these predicted proteins were compared to sequences compiled in GenBank EMBL databases, 20 ORFs showed significant similarity to proteins of bacteriophage 29 that infects *B. subtilis* (Martin et al., 1996). The similar proteins corresponded to those involved in DNA replication (terminal protein and DNA polymerase), structural and morphogenic proteins (major head, collar, connector, tail, and encapsidation proteins), and proteins involved in lysis function (holin and lysozyme). In its strategy of lysis, the holin gene product inserts itself into the cell membrane, allowing access of the lysozyme to the peptidoglycan. Expression of the Cp-1 holin protein in *E. coli* results in cell death after 2-hours of induction, but did not lead to lysis (Garcia et al., 1997). Cells harboring a plasmid construction with holin and lysozyme genes together did lyse after induction and the

viability loss was similar to that of the culture expressing holin alone. Cloning of these lytic genes in *S. pneumoniae* showed that both genes had the same effect as in *E. coli*. That is, holin itself did not lyse the culture but the viability loss was noticeable, whereas both holin and lysozyme together were capable of lysing M31, an amidase
5 deleted mutant (Garcia et al., 1997).

Recently, a small portion (~4 kbp) of a second *S. pneumoniae* phage, Dp-1, has been sequenced (Sheehan et al., 1997). This portion contains the genes coding for the lytic system (Sheehan et al., 1997) and shows a modular organization similar to that described for Cp-1. However, in this case, a single chimeric protein appears to be
10 made in which the N-terminal domain is highly similar to that of the murein hydrolase coded by a gene found in the phage BK5-T that infects *Lactococcus lactis*, and the C-terminal domain is homologous to holins. Thus, both functions appear to have been combined in a novel chimeric protein.

Bacteriophage Dp-1 was obtained from Dr. P. Garcia (Departamento de
15 Microbiologia Molecular, Centro de Departamento de Investigaciones Biologicas, Consejo Superior de Investigaciones Cientificas, Velazquez, Madrid, Spain). We found that Dp-1 has a double-stranded DNA genome of 56,506 bp, predicted to encode 85 ORFs greater than 33 amino acids and with upstream Shine-Dalgarno motifs for translation initiation (Tables 28 & 30, and Fig. 6). Computational analysis
20 of the predicted protein products of *Streptococcus* bacteriophage Dp-1 protein products, which detected homologs in public databases, are listed in Table 31, along with the accompanying list of related proteins.

From this analysis, it is apparent that several predicted genes of Dp-1 encode polypeptides that are related to structural proteins. ORFs 001, 002, 004, and 030 are
25 predicted to encode tail proteins, minor structural proteins, and minor capsid proteins (Table 31). We also note the identification of several gene products that are likely involved in DNA synthesis. These include ORF 3 which encodes DNA polymerase, ORF 8 which encodes a SWI/SNF helicase-related protein, ORF 10 encodes a protein showing homology to recA, and ORF 13 encodes a dnaZX-like ORF.

30 In *E. coli*, RapA encodes an RNA polymerase (RNAP)-associated protein with ATPase activity and which is a homolog of the eukaryotic SWI/SNF family, a set of proteins whose members are involved are involved in transcription activation, nucleosome remodeling, and DNA repair. RapA forms a stable complex with RNAP,

as if it were a subunit of RNAP and it is possible that the ORF 8 product behaves similarly or in a dominant-negative fashion to inhibit the activity of RapA. Mutation of the essential *E. coli* dnaZX results in a block in DNA chain elongation during replication (Maki et al., 1988). The dnaZX gene has only one open reading frame for
 5 a 71-kDa polypeptide from which the two distinct DNA polymerase III holoenzyme subunits, tau (71 kDa) and gamma (47 kDa), are produced. The tau subunit is the precursor of the gamma subunit, and the gamma subunit is produced by a -1 frameshift causing early termination of translation (Tsuchihashi et al., 1990). These proteins show single-strand DNA binding properties that is ATPase (and dATPase)
 10 dependent and are thought to increasing the processivity of the core DNA polymerase enzyme (Lee et al., 1987).

There are several Dp-1 ORFs which encode proteins predicted to play a role in cellular metabolic pathways. These include polypeptides involved in coenzyme PQQ synthesis (ORFs 20, 29, 38). Pyrrolo-quinoline quinone (PQQ) is the non-covalently
 15 bound prosthetic group of many quinoproteins catalysing reactions in the periplasm of Gram-negative bacteria. Most of these involve the oxidation of alcohols or aldose sugars. Interestingly, ORFs 20, 29, and 30 also show homology to the exoenzyme S regulon (Frank, 1997). Proteins encoded by the *P. aeruginosa* exoenzyme S regulon may be involved in a contact-mediated translocation mechanism to transfer anti-host
 20 factors directly into eukaryotic cells disrupting eukaryotic signal transduction through ADP-ribosylation (Frank, 1997).

There is also a protein with similarity to GTP cyclohydrolase I (ORF 21) and ORF 41 which shows homology to dUTPase (Table 31). GTP cyclohydrolase I is an enzyme that catalyzes the first reaction in the pathway for the biosynthesis of the
 25 pteridine, a cofactor of the monooxygenases of the aromatic amino acids. Disruption of the homologous gene in *Saccharomyces cerevisiae* leads to a recessive conditional lethality due to folinic acid auxotrophy, that can be complemented with the mammalian or bacterial GTP cyclohydrolase I enzymes (Nardese et al., 1996; Mancini et al., 1999).

30 ORF 16 shows high homology to autolysin. This region of the phage sequence was previously reported (Sheehan et al., 1997) and encompasses ~ 4 kbp of our sequence. The sequence published by (Sheehan et al., 1997) is shown in Table 32.

Thus, the present invention provides a nucleic acid sequence obtained from *Streptococcus* bacteriophage Dp-1 comprising at least a portion of a phage Dp-1 ORF,
 35 preferably an inhibitory ORF, and more preferably at least a portion of one of the genes described above with anti-microbial activity. For example, ORF 013 encodes a

protein with homology to the gamma subunit of DNA polymerase (*dnaX* gene). This protein may act in a dominant-negative fashion to sequester the host DNA polymerase for its own replication, thus inhibiting host DNA replication. The *dnaX* gene product is essential for *E. coli* replication (Kodaira et al., 1983).

5 In certain preferred embodiments of the present invention, the bacterial target of a bacteriophage inhibitor ORF product, e.g., an inhibitory protein or polypeptide, is encoded by a *Streptococcus* nucleic acid coding sequence from a host bacterium for bacteriophage Dp-1. As above, possible target sequences are described herein by reference to sequence source sites. The sequence encoding the target preferably
10 corresponds to a *Streptococcus* nucleic acid sequence available from The Institute for Genomic Research (TIGR), or available from GenBank or other public database. The TIGR *Streptococcus* sequences are publicly available at The Institute for Genomics Research at URL: <http://www.tigr.org>

The amino acid sequence of a polypeptide target is readily provided by
15 translating the corresponding coding region. For the sake of brevity, the sequences are not reproduced herein. Also, in preferred embodiments, a target sequence corresponds to a *Streptococcus pneumoniae* coding sequences corresponding to a sequence listed in Table 33 herein. Sequences for other Streptococcal species are also available from TIGR and/or from GenBank. The listing in Table 33 describes
20 *Streptococcus* sequences currently deposited in GenBank. Again, for the sake of brevity, the sequences are described by reference to the GenBank entries instead of being written out in full herein. In cases where the TIGR or GenBank entry for a coding region is not complete, the complete sequence can be readily obtained by routine methods, e.g., by isolating a clone in a phage Dp-1 host *Streptococcus* sp.
25 genomic library, and sequencing the clone insert to provide the relevant coding region. The boundaries of the coding region can be identified by conventional sequence analysis and/or by expression in a bacterium in which the endogenous copy of the coding region has been inactivated and using subcloning to identify the functional start and stop codons for the coding region.

30 In the various aspects of this invention involving Dp-1 sequences, preferably the sequence is preferably not contained in the sequence described in Sheehan et al., 1997 (Table 32).

Validating Identified Inhibitory Phage ORFs

35 A fifth step involves validating the identified phage inhibitor ORF by independent methods, and delineating further possible smaller segments of the ORFs

that have inhibitory activity. Several methods exist to validate the role of the identified ORF as an inhibitor ORF.

One example utilizes the creation of a mutant variant of the phage ORF in which the candidate ORF carries a partial or complete loss-of-function mutation that is measurable as compared with the non-mutant ORF. Comparison of the effects of expression of the loss of function mutant with the normal ORF provides confirmation of the identification of an inhibitor ORF where the loss-of-function mutant provides a measurably lower level of inhibition, preferably no inhibition. The loss of function may be conditional, *e.g.*, temperature sensitive.

Once validation of the inhibitor ORF is achieved, a bi-directional deletion analysis can be carried out using the same experimental system to identify the minimal polypeptide segment that has inhibitor activity. This may be carried out by a variety of means, *e.g.*, by exonuclease or PCR methodologies, and is used to determine if a relatively small segment of the ORF (*i.e.*, the product of the ORF) still possesses inhibitory activity when isolated away from its native sequence. If so, a portion of the ORF encoding this "active portion" can be used as a template for the synthesis of novel anti-microbial agents and further allowing derivation of the peptide sequence, *e.g.*, using modified peptides and/or peptidomimetics.

In creation of certain peptidomimetics, the peptide backbone is transformed into a carbon-based hydrophobic structure that can retain inhibitor activity against the bacterium. This is done by standard medicinal chemistry methods, typically monitored by measuring growth inhibition of the various molecules in liquid cultures or on solid medium. These mimetics can also represent lead compounds for the development of novel antibiotics.

Recently, a major effort has been undertaken by the pharmaceutical industry and their biotechnology partners for the sequencing of bacterial pathogen genomes. The rationale is that the systematic sequencing of the genome will identify all of the bacterial proteins and therefore this proteome will be the target for designing novel inhibitor antibiotics. Although systematic, this approach has several major problems. The first is that analysis of primary amino acid sequences of bacterial proteins does not immediately reveal which protein will be essential for viability of the bacterium, and target validation is thus a major issue. The second problem is one of redundancy, as several biochemical pathways are either structurally duplicated in bacteria (different isoforms of the same enzyme), or functionally duplicated by the presence of salvage pathways in the event of a metabolic block in one pathway (different nutritional conditions). The third is that even a valid target may not be structurally or

functionally amenable to inhibition by small molecules because of inaccessibility (sequestration of target).

Therefore, there is considerable interest within the pharmaceutical and biotechnology industry in identifying key targets for drug discovery amongst the mass
5 of novel targets generated by large-scale genomic sequencing projects.

On the other hand, and underscoring the instant invention, the phages herein described have, over millions of years, evolved specific mechanisms to target such key biochemical pathways and proteins. In the few cases where inhibition by phages has been elucidated (*e.g.*, see ref. 3), such bacterial targets are invariably rate-limiting
10 in their respective biochemical pathways, are not redundant, and/or are readily accessible for inhibition by the phage (or by another inhibitory compound). Therefore, the sixth step of this invention involves identifying the host biochemical pathways and proteins that are targeted by the phage inhibitory mechanisms.

15 Identifying, Validating, and Characterizing Bacterial Host Target Proteins and Affected Pathways

A rationale for this step is that the inhibitor ORF product from the phage physically interacts with and/or modifies certain microbial host components to block their function. Exemplary approaches which can be used to identify the host bacterial
20 pathways and proteins that interact with, and preferably also are inhibited by, phage ORF product(s) are described below.

One approach is a genetic screen to determine physiological protein:protein interaction, for example, using a yeast two hybrid system. In this assay, the phage ORF is fused to the carboxyl terminus of the yeast Gal4 activation domain II (amino
25 acids 768-881) to create a bait vector. A cDNA library of cloned *S. aureus* sequences which have been engineered into a plasmid where the *S. aureus* sequences are fused to the DNA binding domain of Gal4 is also generated. These plasmids are introduced alone, or in combination, into yeast strain Y190 - previously engineered with chromosomally integrated copies of the *E. coli lacZ* and the selectable HIS3 genes,
30 both under Gal4 regulation (Durfee, T., Becherer, K., Chen, P.-L., Yeh, S.-H., Yang, Y., Kilburn, A.E., Lee, W.-H., and Elledge, S.J. (1993). *Genes & Dev.* 7, 555-569). If the two proteins expressed in yeast interact, the resulting complex will activate transcription from promoters containing Gal4 binding sites. A *lacZ* and His3 gene, each driven by a promoter containing Gal4 binding sites, have been integrated into the
35 genome of the host yeast system used for measuring protein-protein interactions. Such a system provides a physiological environment in which to detect potential protein interactions. This system has been extensively used to identify novel protein-protein

interaction partners and to map the sites required for interaction (for example, to identify interacting partners of translation factors (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). *Mol & Cell Biology* 18, 2697-2711), transcription factors (Katagiri, T., Saito, H., Shinohara, A., Ogawa, H., Kamada, N., Nakamura, Y., and Miki, Y. (1998). *Genes, Chromosomes & Cancer* 21, 217-222), and proteins involved in signal transduction (Endo, T.A., Masuhara, M., Yokouchi, M., Suzuki, R., Sakamoto, H., Mitsui, K., Matsumoto, A., Tanimura, S., Ohtsubo, M., Misawa, H., Miyazaki, T., Leonor N., Taniguchi, T., Fujita, T., Kanakura, Y., Komiya, S., and Yoshimura, A. *Nature*. 387, 921-924). This approach has also been used in many published reports to identify interaction between mammalian viral and mammalian cell proteins.

For example, the non-structural protein NS1 of parvovirus is essential for viral DNA amplification and gene expression and is also the major cytopathic effector of these viruses. A yeast two-hybrid screen with NS1 identified a novel cellular protein of unknown function that interacts with NS-1, called SGT, for small glutamine-rich tetratricopeptide repeat (TPR)-containing protein (Cziepluch C. Kordes E. Poirey R. Grewenig A. Rommelaere, J, and Jauniaux JC. (1998) *J Virol*. 72, 4149-4156). In another screen, the adenovirus E3 protein was recently shown to interact with a novel tumor necrosis factor alpha-inducible protein and to modulate some of the activities of E3 (Li Y. Kang J. and Horwitz M.S. (1998). *Mol & Cell Biol*. 18, 1601-1610). In yet another recent screen, the herpes simplex virus 1 alpha regulatory protein ICP0 was found to interact with (and stabilize) the cell cycle regulator cyclin D3 (Kawaguchi Y. Van Sant C. and Roizman B. (1997). *J Virol*. 71, 7328-7336).

Another two-hybrid system for identifying protein:protein interactions is commercially available from STRATEGENETM as the CYTO-TRAPTM system (Chang et al., *Strategies Newsletter* 11(3), 65-68 (1998)(from Stratagene)). The system is a yeast-based method for detecting protein:protein interactions *in vivo*, using activation of the Ras signal transduction cascade by localizing a signal pathway component, human Sos (hSos), to its activation site in the yeast plasma membrane. The system uses a temperature-sensitive *Saccharomyces cerevisiae* mutant, strain cdc25H, which contains a point mutation at amino acid residue 1328 of the cdc25 gene. This gene encodes a guanyl nucleotide exchange factor which binds and activates Ras, leading to cell growth. The mutation in the cdc25 gene prevents host growth at 37°C, but at a permissive temperature of 25°C, growth is normal. The system utilizes the ability of (hSos) to complement the cdc25 defect and activate the yeast Ras signaling pathway. Once (hSos) is expressed and localized to the plasma membrane, the cdc25H yeast strain grows at 37°C. Localizing hSos to the plasma

membrane occurs through a protein:protein interaction. A protein of interest, or bait, is expressed as a fusion protein with hSos. The library, or target proteins are expressed with the myristylation membrane-localization signal. The yeast cells are then incubated under restrictive conditions (37°C). If the bait and the target protein interact, the hSos protein is recruited to the membrane, activating the Ras signaling pathway and allowing the cdc25H yeast strain to grow at the restrictive temperature.

The protein targets of phage inhibitory ORFs can also be identified using bacterial genetic screens. One approach involves the overexpression of a phage inhibitory protein in mutagenized bacterial host species, followed by plating the cells and searching for colonies that can survive the antimicrobial activity of the inhibitory ORF. These colonies are then grown, their DNA extracted, and cloned into an expression vector that contains a replicon of a different incompatibility group from the plasmid expressing the original ORF. This library is then introduced into a wild-type host bacterium in conjunction with an expression vector driving synthesis of the phage ORF, followed by selection for surviving bacteria. Thus, bacterial DNA fragments from the survivors presumably contain a DNA fragment from the original mutagenized host bacterial genome that can protect the cell from the antimicrobial activity of the inhibitory phage ORF. This fragment can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach enables one to determine the targets and pathways that are affected by the killing function.

A second approach is based on identifying protein:protein interactions between the phage ORF product and bacterial *S. aureus*, e.g., proteins using a biochemical approach based, for example, on affinity chromatography. This approach has been used, for example, to identify interactions between lambda phage proteins and proteins from their *E. coli* host (Sopta, M., Carthew, R.W., and Greenblatt, J. (1985) *J. Biol. Chem.* 260, 10353-10369). The phage ORF is fused to a peptide tag (e.g. glutathione-S-transferase ("GST"), 6xHIS, ("HIS") and/or calmodulin binding protein ("CPB")) within a commercially available plasmid vector that directs high level expression on induction of a suitably responsive promoter driving the fusion's expression. The translated fusion protein is expressed in *E. coli*, purified, and immobilized on a solid phase matrix via, for example the tag. Total cell extracts from the host bacterium, e.g., *S. aureus*, are then passed through the affinity matrix containing the immobilized phage ORF fusion protein; host proteins retained on the column are then eluted under different conditions of ionic strength, pH, detergents etc., and characterized by gel electrophoresis and other techniques. Appropriate controls are run to guard against nonspecific binding to the resin. Target proteins thus

recovered should be enriched for the phage protein/peptide of interest and are subsequently electrophoretically or otherwise separated, purified, sequenced, or biochemically analyzed. Usually sequencing entails individual digestion of the proteins to completion with a protease (*e.g.*-trypsin), followed by molecular mass and amino acid composition and sequence determination using, for example, mass spectrometry, *e.g.*, by MALDI-TOF technology (Qin, J., Fenyo, D., Zhao, Y., Hall, W.W., Chao, D.M., Wilson, C.J., Young, R.A. and Chait, B.T. (1997). *Anal. Chem.* 69, 3995-4001).

The sequence of the individual peptides from a single protein are then analyzed by the bioinformatics approach described above to identify the *S. aureus* protein interacting with the phage ORF. This analysis is performed by a computer search of the *S. aureus* genome for an identified sequence. Alternatively, all tryptic peptide fragments of the *S. aureus* genome can be predicted by computer software, and the molecular mass of such fragments compared to the molecular mass of the peptides obtained from each interacting protein eluted from the affinity matrix. The responsible gene sequence can be obtained, for example by using synthetic degenerate nucleic acid sequences to pull out the corresponding homologous bacterial sequence. Alternatively, antibodies can be generated against the peptide and used to isolate nascent peptide/mRNA transcript complexes, from which the mRNA can be reverse transcribed, cloned, and further characterized using the procedures discussed herein.

A variety of other binding assay methods are known in the art and can be used to identify interactions between phage proteins and bacterial proteins or other bacterial cell components. Such methods that allow or provide identification of the bacterial component can be used in this invention for identifying putative targets.

Validation of the interaction between the phage ORF product and the bacterial proteins or other components can be obtained by a second independent assay (*e.g.*, co-immunoprecipitation or protein-protein crosslinking experiments (Qiu, H., Garcia-Barrio, M.T., and Hinnebusch, A.G. (1998). *Mol & Cell Biology* 18, 2697-2711; Brown, S. and Blumenthal, T. (1976). *Proc. Natl. Acad. Sci. USA* 73, 1131-1135)).

Finally, the essential nature of the identified bacterial proteins is preferably determined genetically by creating a constitutive or inducible partial or complete loss-of-function mutation in the gene encoding the identified interacting bacterial protein. This mutant is then tested for bacterial survival and replication.

The protein target of the phage inhibitor function can also be identified using a genetic approach. Two exemplary approaches will be delineated here. The first approach involves the overexpression of a predetermined phage inhibitor protein in mutagenized host bacteria, *e.g.*, *S. aureus*, followed by plating the cells and searching

for colonies that can survive the inhibitor. These colonies will then be grown, their DNA extracted and cloned into an expression vector that contains a replicon of a different incompatibility group, and preferably having a different selectable marker than the plasmid expressing the phage inhibitor. Thus, host DNA fragments from the mutant that can protect the cell from phage ORF inhibition can be sequenced and compared with that of the bacterial host to determine in which gene the mutation lies. This approach allows rapid determination of the targets and pathways that are affected by the inhibitor.

Alternatively, the bacterial targets can be determined in the absence of selecting for mutations using an approach known as "multicopy suppression". In this approach, the DNA from the wild type host is cloned into an expression vector that can coexist, as previously described, with one containing a predetermined phage inhibitor. Those plasmids that contain host DNA fragments and genes that protect the host from the phage inhibitor can then be isolated and sequenced to identify putative targets and pathways in the host bacteria.

Regardless of the specific mode of identification, screening assays may additionally utilize gene fusions to specific "reporter genes" to identify a bacterial gene(s) whose expression is affected when the host target pathway is affected by the phage inhibitor. Such gene fusions can be used to search a number of small molecule compounds for inhibitors that may affect this pathway and thus cause cell inhibition. This approach will allow the screening of a large number of molecules on petri dishes or 96-well format by monitoring for a simple color change in the bacterial colonies. In this manner, we can validate host targets and classes of compounds for further study and clinical development. These inhibitors also represent lead compounds for the development of other antibiotics.

Bioinformatics and comparative genomics are preferably then applied to the identified bacterial gene products to predict biochemical function. The biochemical activity of the protein can be verified *in vitro* in cell free assays or *in vivo* in intact cells. *In vitro* biochemical assays utilizing cell-free extracts or purified protein are established as a basis for the screening and development of inhibitors.

These inhibitors, preferably small molecule inhibitors, may comprise peptides, antibodies, products from natural sources such as fungal or plant extracts or small molecule organic compounds. In general, small molecule organic compounds are preferred. These compounds may, for example, be identified within large compound libraries, including combinatorial libraries. For example, a plurality of compounds, preferably a large number of compounds can be screened to determine whether any of the compounds binds or otherwise disrupts or inhibits the identified bacterial target.

Compounds identified as having any of these activities can then be evaluated further in cell culture and/or animal model systems to determine the pharmacological properties of the compound, including the specific anti-microbial ability of the compound.

- 5 For mixtures of natural products, including crude preparations, once a preparation or fraction of a preparation is shown to have an anti-microbial activity, the active substance can be isolated and identified using techniques well known in the art, if the compound is not already available in a purified form.

- 10 Identified compounds possessing anti-microbial activity and similar compounds having structural similarity can be further evaluated and, if necessary, derivatized according to synthesis and/or modification methods available in the art selected as appropriate for the particular starting molecule.

Derivatization of identified anti-microbials

- 15 In cases where the identified anti-microbials above might represent peptidal compounds, the *in vivo* effectiveness of such compounds may be advantageously enhanced by chemical modification using the natural polypeptide as a starting point and incorporating changes that provide advantages for use, for example, increased stability to proteolytic degradation, reduced antigenicity, improved tissue penetration,
20 and/or improved delivery characteristics.

- In addition to active modifications and derivative creations, it can also be useful to provide inactive modifications or derivatives for use as negative controls or introduction of immunologic tolerance. For example, a biologically inactive derivative which has essentially the same epitopes as the corresponding natural
25 antimicrobial can be used to induce immunological tolerance in a patient being treated. The induction of tolerance can then allow uninterrupted treatment with the active anti-microbial to continue for a significantly longer period of time.

- Modified anti-microbial polypeptides and derivatives can be produced using a number of different types of modifications to the amino acid chain. Many such
30 methods are known to those skilled in the art. The changes can include, for example, reduction of the size of the molecule, and/or the modification of the amino acid sequence of the molecule. In addition, a variety of different chemical modifications of the naturally occurring polypeptide can be used, either with or without modifications to the amino acid sequence or size of the molecule. Such chemical modifications can,
35 for example, include the incorporation of modified or non-natural amino acids or non-amino acid moieties during synthesis of the peptide chain, or the post-synthesis modification of incorporated chain moieties.

The oligopeptides of this invention can be synthesized chemically or through an appropriate gene expression system. Synthetic peptides can include both naturally occurring amino acids and laboratory synthesized, modified amino acids.

Also provided herein are functional derivatives of anti-microbial proteins or polypeptides. By "functional derivative" is meant a "chemical derivative,"
5 "fragment," "variant," "chimera," or "hybrid" of the polypeptide or protein, which terms are defined below. A functional derivative retains at least a portion of the function of the protein, for example reactivity with a specific antibody, enzymatic activity or binding activity.

10 A "chemical derivative" of the complex contains additional chemical moieties not normally a part of the protein or peptide. Such moieties may improve the molecule's solubility, absorption, biological half-life, and the like. The moieties may alternatively decrease the toxicity of the molecule, eliminate or attenuate any
15 such effects are disclosed in Alfonso and Gennaro (1995). Procedures for coupling such moieties to a molecule are well known in the art. Covalent modifications of the protein or peptides are included within the scope of this invention. Such modifications may be introduced into the molecule by reacting targeted amino acid residues of the peptide with an organic derivatizing agent that is capable of reacting
20 with selected side chains or terminal residues, as described below.

Cysteiny l residues most commonly are reacted with alpha-haloacetates (and corresponding amines), such as chloroacetic acid or chloroacetamide, to give carboxymethyl or carboxyamidomethyl derivatives. Cysteiny l residues also are derivatized by reaction with bromotrifluoroacetone, chloroacetyl phosphate, N-
25 alkylmaleimides, 3-nitro-2-pyridyl disulfide, methyl 2-pyridyl disulfide, p-chloro-mercuribenzoate, 2-chloromercuri-4-nitrophenol, or chloro-7-nitrobenzo-2-oxa-1,3-diazole.

Histidyl residues are derivatized by reaction with diethylprocarbonate at pH 5.5-7.0 because this agent is relatively specific for the histidyl side chain. Para-
30 bromophenacyl bromide also is useful; the reaction is preferably performed in 0.1 M sodium cacodylate at pH 6.0.

Lysiny l and amino terminal residues are reacted with succinic or other carboxylic acid anhydrides. Derivatization with these agents has the effect of reversing the charge of the lysiny l residues. Other suitable reagents for derivatizing
35 primary amine- containing residues include imidoesters such as methyl picolinimate; pyridoxal phosphate; pyridoxal; chloroborohydride;

trinitrobenzenesulfonic acid; O-methylisourea; 2,4 pentanedione; and transaminase-catalyzed reaction with glyoxylate.

Arginyl residues are modified by reaction with one or several conventional reagents, among them phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, and ninhydrin. Derivatization of arginine residues requires that the reaction be performed in alkaline conditions because of the high pK_a of the guanidine functional group. Furthermore, these reagents may react with the groups of lysine as well as the arginine alpha-amino group.

Tyrosyl residues are well-known targets of modification for introduction of spectral labels by reaction with aromatic diazonium compounds or tetranitromethane. Most commonly, N-acetylimidizol and tetranitromethane are used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively.

Carboxyl side groups (aspartyl or glutamyl) are selectively modified by reaction carbodiimide ($R'-N-C-N-R'$) such as 1-cyclohexyl-3-(2-morpholinyl(4-ethyl) carbodiimide or 1-ethyl-3-(4-azonia-4,4-dimethylpentyl) carbodiimide. Furthermore, aspartyl and glutamyl residues are converted to asparaginy and glutaminy residues by reaction with ammonium ions.

Glutaminy and asparaginy residues are frequently deamidated to the corresponding glutamyl and aspartyl residues. Alternatively, these residues are deamidated under mildly acidic conditions. Either form of these residues falls within the scope of this invention.

Derivatization with bifunctional agents is useful, for example, for cross-linking component peptides to each other or the complex to a water-insoluble support matrix or to other macromolecular carriers. Commonly used cross-linking agents include, for example, 1,1-bis (diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, for example, esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), and bifunctional maleimides such as bis-N-maleimido-1,8-octane. Derivatizing agents such as methyl-3-[p-azidophenyl] dithiolpropioimide yield photoactivatable intermediates that are capable of forming crosslinks in the presence of light. Alternatively, reactive water-insoluble matrices such as cyanogen bromide-activated carbohydrates and the reactive substrates described in U.S. Patent Nos. 3,969,287; 3,691,016; 4,195,128; 4,247,642; 4,229,537; and 4,330,440 are employed for protein immobilization.

Other modifications include hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of seryl or threonyl residues, methylation of the alpha-amino groups of lysine, arginine, and histidine side chains (Creighton, T.E.,

Proteins: Structure and Molecular Properties, W.H. Freeman & Co., San Francisco, pp. 79-86 (1983)), acetylation of the N-terminal amine, and, in some instances, amidation of the C-terminal carboxyl groups.

Such derivatized moieties may improve the stability, solubility, absorption,
5 biological half life, and the like. The moieties may alternatively eliminate or attenuate any undesirable side effect of the protein complex. Moieties capable of mediating such effects are disclosed, for example, in Alfonso and Gennaro (1995).

The term "fragment" is used to indicate a polypeptide derived from the amino acid sequence of the protein or polypeptide having a length less than the full-length
10 polypeptide from which it has been derived. Such a fragment may, for example, be produced by proteolytic cleavage of the full-length protein. Preferably, the fragment is obtained recombinantly by appropriately modifying the DNA sequence encoding the proteins to delete one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

15 Another functional derivative intended to be within the scope of the present invention is a "variant" polypeptide that either lacks one or more amino acids or contains additional or substituted amino acids relative to the native polypeptide. The variant may be derived from a naturally occurring polypeptide by appropriately modifying the protein DNA coding sequence to add, remove, and/or to modify codons
20 for one or more amino acids at one or more sites of the C-terminus, N-terminus, and/or within the native sequence.

A functional derivative of a protein or polypeptide with deleted, inserted and/or substituted amino acid residues may be prepared using standard techniques well-known to those of ordinary skill in the art. For example, the modified
25 components of the functional derivatives may be produced using site-directed mutagenesis techniques (as exemplified by Adelman et al., 1983, *DNA* 2:183; Sambrook et al., 1989) wherein nucleotides in the DNA coding sequence are modified such that a modified coding sequence is produced, and thereafter expressing this recombinant DNA in a prokaryotic or eukaryotic host cell, using techniques such as
30 those described above. Alternatively, components of functional derivatives of complexes with amino acid deletions, insertions and/or substitutions may be conveniently prepared by direct chemical synthesis, using methods well-known in the art.

Insofar as other anti-microbial inhibitor compounds identified by the invention
35 described herein may not be peptidal in nature, other chemical techniques exist to allow their suitable modification, as well, and according the desirable principles discussed above.

Administration and Pharmaceutical Compositions

For the therapeutic and prophylactic treatment of infection, the preferred method of preparation or administration of anti-microbial compounds will generally vary depending on the precise identity and nature of the anti-microbial being delivered. Thus, those skilled in the art will understand that administration methods known in the art will also be appropriate for the compounds of this invention.

The particularly desired anti-microbial can be administered to a patient either by itself, or in pharmaceutical compositions where it is mixed with suitable carriers or excipient(s). In treating an infection, a therapeutically effective amount of an agent or agents is administered. A therapeutically effective dose refers to that amount of the compound that results in amelioration of one or more symptoms of bacterial infection and/or a prolongation of patient survival or patient comfort.

Toxicity, therapeutic and prophylactic efficacy of anti-microbials can be determined by standard pharmaceutical procedures in cell cultures and/or experimental organisms such as animals, *e.g.*, for determining the LD_{50} (the dose lethal to 50% of the population) and the ED_{50} (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio LD_{50}/ED_{50} . Compounds that exhibit large therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED_{50} with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized.

For any compound identified and used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. Such information can be used to more accurately determine useful doses in organisms such as plants and animals, preferably mammals, and most preferably humans. Levels in plasma may be measured, for example, by HPLC or other means appropriate for detection of the particular compound.

The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition (see *e.g.* Fingl et. al., in *The Pharmacological Basis of Therapeutics*, 1975, Ch. 1 p.1).

It should be noted that the attending physician would know how and when to terminate, interrupt, or adjust administration due to toxicity, organ dysfunction, or other systemic malady. Conversely, the attending physician would also know to adjust treatment to higher levels if the clinical response were not adequate (precluding

toxicity). The magnitude of an administered dose in the management of the disorder of interest will vary with the severity of the condition to be treated and the route of administration. The severity of the condition may, for example, be evaluated, in part, by standard prognostic evaluation methods. Further, the dose and perhaps dose
5 frequency, will also vary according to the age, body weight, and response of the individual patient. A program comparable to that discussed above also may be used in veterinary or phyto medicine.

Depending on the specific infection target being treated and the method selected, such agents may be formulated and administered systemically or locally, i.e.,
10 topically. Techniques for formulation and administration may be found in Alfonso and Gennaro (1995). Suitable routes may include, for example, oral, rectal, transdermal, vaginal, transmucosal, intestinal, parenteral, intramuscular, subcutaneous, or intramedullary injections, as well as intrathecal, intravenous, or intraperitoneal injections.

15 For injection, the agents of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks' solution, Ringer's solution, or physiological saline buffer. For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art.

20 Use of pharmaceutically acceptable carriers to formulate identified anti-microbials of the present invention into dosages suitable for systemic administration is within the scope of the invention. With proper choice of carrier and suitable manufacturing practice, the compositions of the present invention, in particular those formulated as solutions, may be administered parenterally, such as by intravenous
25 injection. Appropriate compounds can be formulated readily using pharmaceutically acceptable carriers well known in the art into dosages suitable for oral administration. Such carriers enable the compounds of the invention to be formulated as tablets, pills, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated.

30 Agents intended to be administered intracellularly may be administered using techniques well known to those of ordinary skill in the art. For example, such agents may be encapsulated into liposomes, then administered as described above. Liposomes are spherical lipid bilayers with aqueous interiors. All molecules present in an aqueous solution at the time of liposome formation are incorporated into the
35 aqueous interior. The liposomal contents are both protected from the external microenvironment and, because liposomes fuse with cell membranes, are efficiently

delivered into the cell cytoplasm. Additionally, due to their hydrophobicity, small organic molecules may be directly administered intracellularly.

Pharmaceutical compositions suitable for use in the present invention include compositions wherein the active ingredients are contained in an effective amount to achieve the intended purpose. Determination of the effective amounts is well within the capability of those skilled in the art.

In addition to the active ingredients, these pharmaceutical compositions may contain suitable pharmaceutically acceptable carriers comprising excipients and auxiliaries which facilitate processing of the active compounds into preparations which can be used pharmaceutically. The preparations formulated for oral administration may be in the form of tablets, dragees, capsules, or solutions, including those formulated for delayed release or only to be released when the pharmaceutical reaches the small or large intestine.

The pharmaceutical compositions of the present invention may be manufactured in a manner that is itself known, *e.g.*, by means of conventional mixing, dissolving, granulating, dragee-making, levitating, emulsifying, encapsulating, entrapping or lyophilizing processes.

Pharmaceutical formulations for parenteral administration include aqueous solutions of the active anti-microbial compounds in water-soluble form. Alternatively, suspensions of the active compounds may be prepared as appropriate oily injection suspensions. Suitable lipophilic solvents or vehicles include fatty oils such as sesame oil, or synthetic fatty acid esters, such as ethyl oleate or triglycerides, or liposomes. Aqueous injection suspensions may contain substances which increase the viscosity of the suspension, such as sodium carboxymethyl cellulose, sorbitol, or dextran. Optionally, the suspension may also contain suitable stabilizers or agents which increase the solubility of the compounds to allow for the preparation of highly concentrated solutions.

Pharmaceutical preparations for oral use can be obtained by combining the active compounds with solid excipient, optionally grinding a resulting mixture, and processing the mixture of granules, after adding suitable auxiliaries, if desired, to obtain tablets or dragee cores. Suitable excipients are, in particular, fillers such as sugars, including lactose, sucrose, mannitol, or sorbitol; cellulose preparations such as, for example, maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP). If desired, disintegrating agents may be added, such as the cross-linked polyvinyl pyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

Dragee cores are provided with suitable coatings. For this purpose, concentrated sugar solutions may be used, which may optionally contain gum arabic, talc, polyvinyl pyrrolidone, carbopol gel, polyethylene glycol, and/or titanium dioxide, lacquer solutions, and suitable organic solvents or solvent mixtures.

- 5 Dyestuffs or pigments may be added to the tablets or dragee coatings for identification or to characterize different combinations of active compound doses.

Pharmaceutical preparations which can be used orally include push-fit capsules made of gelatin, as well as soft, sealed capsules made of gelatin and a plasticizer, such as glycerol or sorbitol. The push-fit capsules can contain the active
10 ingredients in admixture with filler such as lactose, binders such as starches, and/or lubricants such as talc or magnesium stearate and, optionally, stabilizers. In soft capsules, the active compounds may be dissolved or suspended in suitable liquids, such as fatty oils, liquid paraffin, or liquid polyethylene glycols. In addition, stabilizers may be added.

- 15 The above methodologies may be employed either actively or prophylactically against an infection of interest.

Computer-related Aspects and Embodiments

In addition to the provision of compounds as chemical entities, nucleotide
20 sequences, or fragments thereof at least 95%, preferably at least 97%, more preferably at least 99%, and most preferably at least 99.9% identical to phage inhibitor sequences can also be provided in a variety of additional media to facilitate various uses.

Thus, as used in this section, "provided" refers to an article of manufacture, rather than an actual nucleic acid molecule, which contains a nucleotide sequence of
25 the present invention; e.g., a nucleotide sequence of an exemplary bacteriophage or a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of an unsequenced phage listed in Table 1, preferably of bacteriophage
30 77 (*S. aureus* host) or bacteriophage 3A (*S. aureus* host) or bacteriophage 96 (*S. aureus* host). Such an article provides a large portion of the particular bacteriophage genome or bacterial gene and parts thereof (e.g., a bacteriophage open reading frame (ORF)) in a form which allows a skilled artisan to examine and/or analyze the sequence using means not directly applicable to examining the actual genome or gene,
35 or subset thereof as it exists in nature or in purified form as a chemical entity.

In one application of this aspect, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer

readable media" refers to any medium that can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories, such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create an article of manufacture which includes one or more computer readable media having recorded thereon a nucleotide sequence or sequences of the present invention. Likewise, it will be clear to those of skill how additional computer readable media that may be developed also can be used to create analogous manufactures having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently known methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can, for example, be presented in a word processing test file, formatted in commercially available software such as WordPerfect and Microsoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase, Oracle, or the like. A skilled artisan can readily adapt any number of data processor structuring formats (e.g., text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. Thus, by providing in computer readable form a nucleotide sequence of an unsequenced bacteriophage, such as an exemplary bacteriophage listed in Table 1 or of a sequence encoding a bacterial target or a fragment thereof, preferably a nucleotide sequence at least 95%, more preferably at least 99% and most preferably at least 99.9% identical to such a bacteriophage or bacterial sequence, for example, to a polynucleotide of bacteriophage 77 (*S. aureus* host) or bacteriophage 3A (*S. aureus* host) bacteriophage

96 (*S. aureus* host), bacteriophage 44AHJD (*S. aureus* host), bacteriophage Dp-1 (*Streptococcus pneumoniae* host), or bacteriophage 182 (*Enterococcus* host) the present invention enables the skilled artisan to routinely access the provided sequence information for a wide variety of purposes.

5 Those skilled in the art understand that software can implement a variety of different search or analysis software which implement sequence search and analysis algorithms, *e.g.*, the BLAST (Altschul et al., J. Mol. Biol. 215:403410 (1990) and BLAZE (Brutlag et al., Comp. Chem 17:203-207 (1993)) search algorithms. For example, such search algorithms can be implemented on a Sybase system and used to
10 identify open reading frames (ORFs) within the bacteriophage genome which contain homology to ORFs or proteins from other viruses, *e.g.*, other bacteriophage, and other organisms, *e.g.*, the host bacterium. Among the ORFs discussed herein are protein encoding fragments of the bacteriophage genomes which encode bacteria-inhibiting proteins or fragments.

15 The present invention further provides systems, particularly computer-based systems, which contain the sequence information described. Such systems are designed to identify, among other things, useful fragments of the bacteriophage genomes.

 As used herein, "a computer-based system" refers to the hardware, software,
20 and data storage media used to analyze the nucleotide sequence information of the present invention. The minimum hardware of the computer-based systems of the present invention comprises a central processing unit (CPU), input device, output device, and data storage medium or media. A skilled artisan will readily recognize that any of the currently available general purpose computer-based system are suitable
25 for use in the present invention, as well as a variety of different specialized or dedicated computer-based systems.

 As stated above, the computer-based systems of the present invention comprise data storage media having stored therein a nucleotide sequence of the present invention and the necessary hardware and software for supporting and
30 implementing a search and/or analysis program.

 As used herein, "data storage media" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

35 As used herein, "search program" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means.

Search means are used to identify fragments or regions of the present genomic sequences which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, MacPattern (EMBL), BLASTN and BLASTX (NCBIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches and/or sequence analyses can be adapted for use in the present computer-based systems.

As used herein in connection with sequence searches and analyses, a "target sequence" can be any DNA or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. Also, the target sequence length is preferably selected to include sequence corresponding to a biologically relevant portion of an encoded product, for example a region which is expected to be conserved across a range of source organisms. Preferably the sequence length of a target polypeptide sequence is from 5-100 amino acids, more preferably 7-50 or 7-100 amino acids, and still more preferably 10-80 or 10-100 amino acids. Preferably the sequence length of a target polynucleotide sequence is from 15-300 nucleotide residues, more preferably from 21-240 or 21-300, and still more preferably 30-150 or 30-300 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length. Likewise, it may be desirable to search and/or analyze longer sequences.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzymatic active sites and signal sequences. Nucleic acid target motifs include, but are not limited to promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

A variety of structural formats for the input and output devices can be used to input and output the information in the computer-based systems of the present invention. A preferred format for an output device ranks fragments of the bacteriophage or bacterial sequences possessing varying degrees of homology to the

target sequence or target motif. Such presentation provides a skilled artisan with a ranking of sequences which contain various amounts of the target sequence or target motif and identifies the degree of homology contained in the identified fragment.

A variety of comparing methods and/or devices and/or formats can be used to
5 compare a target sequence or target motif with the sequence stored in data storage media to identify sequence fragments of the bacteriophage or bacterium in question. One skilled in the art can readily recognize that any one of the publicly available homology search programs can be used as the search program for the computer-based systems of the present invention. Of course, suitable proprietary systems that may be
10 known to those of skill, or later developed, also may be employed in this regard.

Figure 6 provides a block diagram of a computer system illustrative of embodiments of this aspect of present invention. The computer system 102 includes a processor 106 connected to a bus 104. Also connected to the bus 104 are a main memory 108 (preferably implemented as random access memory, RAM) and a variety
15 of secondary storage devices 110, such as a hard drive 112 and a removable medium storage device 114. The removable medium storage device 114 may represent, for example, a floppy disk drive, a CD-ROM drive, a magnetic tape drive, etc. A removable storage medium 116 (such as a floppy disk, a compact disk, a magnetic tape, etc.) containing control logic and/or data recorded therein may be inserted into
20 the removable medium storage device 114. The computer system 102 includes appropriate software for reading the control logic and/or the data from the removable medium storage device 114, once it is inserted into the removable medium storage device 114.

A nucleotide sequence of the present invention may be stored in a well-known
25 manner in the main memory 108, any of the secondary storage devices 110, and/or a removable storage medium 116. During execution, software for accessing and processing the sequence (such as search tools, comparing tools, etc.) reside in main memory 108, in accordance with the requirements and operating parameters of the operating system, the hardware system and the software program or programs.

30 The data storage medium in which the sequence is embodied and the central processor need not be part of a single stand-alone computer, but may be separated so long as data transfer can occur. For example, the processor or processors being utilized for a search or analysis can be part of one general purpose computer, and the data storage medium can be part of a second general purpose computer connected to a
35 network, or the data storage medium can be part of a network server. As another example the data storage medium can be part of a computer system or network accessible over telephone lines or other remote connection method.

EXAMPLES

Example 1. Growth of *Staph A* bacteriophage 77 and purification of genomic DNA.

5 The *Staphylococcus aureus* propagating strain (PS 77; ATCC #27699) was used as a host to propagate its respective phage 77 (ATCC # 27699-B1). Two rounds of plaque purification of phage 77 were performed on soft agar essentially as described in Sambrook et al (1989). Briefly, the PS 77 strain was grown overnight at 37°C in Nutrient broth [NB: 0.3% Bacto beef extract, 0.5% Bacto peptone (Difco
10 Laboratories) and 0.5% NaCl (w/v)]. The culture was then diluted 20x in NB and incubated at 37°C until the $OD_{540} = .2$ (early log phase) with constant agitation. In order to obtain single plaques, phage 77 was subjected to 10-fold serial dilutions using phage buffer (1 mM $MgSO_4$, 5 mM $MgCl_2$, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 µl of each dilution was used to infect 0.5 ml of the cell suspension in the presence
15 of 400 µg/ml $CaCl_2$. After incubation of 15 min at room temperature (RT), 2 ml of melted soft agar kept at 45°C (NB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm nutrient agar plates (0.3% Bacto Beef extract, 0.5% Bacto peptone, 0.5% NaCl and 1.5% Bacto agar (w/v)). After overnight incubation at 30°C, a single plaque was isolated, resuspended in 1 ml of phage buffer
20 by end over end rotation for 2 hrs at 20°C, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 30°C, a single plaque was isolated and used as a stock.

 The propagation procedure for bacteriophage 77 was modified from the agar layer method of Swanstörn and Adams (1951). Briefly, the PS 77 strain was grown to
25 stationary phase overnight at 37°C in Nutrient broth. The culture was then diluted twenty-fold in NB and incubated at 37°C until the $OD_{540} = .2$. The suspension (15×10^7 Bacteria) was then mixed with 15×10^5 plaque forming units (pfu) to give a ratio of 100-bacteria/phage particle in the presence of 400 µg/ml of $CaCl_2$. After incubation for 15 min at 20°C, 7.5 ml of melted soft agar (NB plus 0.6% agar) were added to the
30 mixture and poured onto the surface of 150 mm nutrient agar plates and incubated 16 hrs at 30°C. To collect the phage plate lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 RPM (2,830xg) in a JA-10 rotor--
35 (Beckman) and the supernatant fluid (lysate) was collected and subjected to a treatment with 10 µg /ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) PEG 8000 and

0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin). The phage suspension was
5 extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS 55 rotor centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000xg) for 24 h at
10 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 mg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of
15 phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris pH 8.0, 1mM EDTA).

Example 2. DNA sequencing of Bacteriophage 77 genome

Four micrograms of phage 77 DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed
20 (550 Sonic Dismembrator™, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0])
25 as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris (pH 8.5).

The ends of the sonicated DNA fragments were repaired with a combination of
30 T4 DNA polymerase and the Klenow fragment of E. coli DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5
35 units of Klenow large fragment (New England Biolabs) for 15 min at room-temperature. The reaction was stopped by two phenol/chloroform extractions and the

DNA was precipitated with ethanol and the final DNA pellet was resuspended in 20 μ l of H₂O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs)-treated pKS II+ vector (Stratagene). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 μ l of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 μ l containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10 β according to standard procedures (Sambrook et al., 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μ l LB and 100 μ g/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS II+ vector. PCR amplification of foreign insert was performed in a 15 μ l reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 μ M primer, 187.5 μ M each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 57°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems). To ensure co-linearity of the sequence data and the genome, all regions of phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye™ terminator cycle sequencing ready reaction kit.

Example 3. Bioinformatic management of primary nucleotide sequence from
Phage 77.

Phage 77 sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of

the contigs. Phage DNA was used directly as sequencing template employing ABI prism BIG DYE™ terminator cycle sequencing ready reaction kit. The complete sequence of bacteriophage 77 is shown in Table 2.

A software program was developed and used on the assembled sequence of bacteriophage 77 to identify all putative ORFs larger than 33 codons. Other ORF identification software can also be utilized, preferably programs which allow alternative start codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI (<http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c>) for the bacterial genetic code.

When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons (start and stop codons) is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence.

Sequence homology (BLAST) searches for each ORF are then carried out using an implementation of BLAST programs, although any of a variety of different sequence comparison and matching programs can be utilized as known to those skilled in the art. Downloaded public databases used for sequence analysis include:

- i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- 30 v) *S. aureus* NCTC 8325 (<ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa>);
- vi) *Streptococcus pyogenes* (<ftp://ftp.genome.ou.edu/pub/strep/strep-1k.fa>);
- vii) *Streptococcus pneumoniae* (ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- viii) *Mycobacterium tuberculosis* CSU#9 (ftp://ftp.tigr.org/pub/data/m_tuberculosis/TB_091097.Z) and
- 35 ix) *Pseudomonas aeruginosa* (<http://www.genome.washington.edu/pseudo/data.html>).

The results of the homology searches performed on the ORFs is shown in Table 5.

Example 4. Subcloning of Bacteriophage 77 ORFs into a Staph A inducible expression system.

The shuttle vector pT0021, in which the firefly luciferase (*lucFF*) expression is controlled by the *ars* (arsenite) promoter/operator (Tauriainen et al., 1997), was modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the hemagglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with *Bam*HI, *Sal*I and *Hind*III cloning sites) is:

5'-gatcccggtcgaccaagcttTACCCATACGACGTCCCAGACTACGCCAGCTGA-3' (where upper case letters denote the nucleotide sequence of the HA tag); the antisense strand HA tag sequence (with a *Hind*III cloning site) is:

5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAAagcttggtcgaccgg-3' (where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with *Bam*HI and *Hind*III. This manipulation resulted in replacement of the *lucFF* gene by the HA tag. This modified shuttle vector containing the *arsenite* inducible promoter, the *arsR* gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A.

Each ORF, encoded by Bacteriophage 77, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon was selected for functional analysis for bacterial inhibition. In total, 98 ORFs were selected and screened as detailed below. A list of these is presented in Table 3. Each individual ORF, from initiation codon to last codon (excluding the stop codon), was amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a *Bam*HI restriction site (5'-cgggatcc-3') and each antisense oligonucleotide targets the penultimate codon (the one before the stop codon) of the ORF and is preceded by a *Sal*I restriction site (5'-gcgtcgaccg-3'). The PCR product of each ORF was gel purified and digested with *Bam*HI and *Sal*I. The digested PCR product was then gel purified using the Qiagen kit as described, ligated into *Bam*HI and *Sal*I digested pTHA vector, and used to transform *E. coli* bacterial strain DH10 β (as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones were picked and their insert sizes were confirmed by PCR analysis

using primers flanking the cloning site. The names and sequences of the primers that were used for the PCR amplification were: HAF:

'TATTATCCAAAACCTTGAACA'; HAR: 'CGGTGGTATATCCAGTGATT'. The sequence integrity of cloned ORFs was verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence could not be achieved by one pass with the sequencing primers, additional internal primers were selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiwirth et al., 1983) was used as a recipient for the expression of recombinant plasmids. Electroporation was performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones was performed on Luria-Broth agar (LB-agar) plates containing 30 µg/ml of kanamycin.

For each ORF introduced in the pTHA plasmid, 3 independent transformants were isolated and used to individually inoculate cultures in 5 ml of TSB containing 30 µg/ml kanamycin, followed by growth to saturation (16 hrs at 30°C). An aliquot of this stationary phase culture was used to generate a frozen glycerol stock of the transformant (stored at -80°C). The remaining culture was used for plasmid DNA extraction. Bacterial cells were harvested by centrifugation at 3000 x g at 22°C for 5 min. The pellet was resuspended in 200 µl 25% sucrose containing 25U/ml of lysostaphin and incubated for 15 min at 37°C. Then, 400 µl of alkaline SDS solution (3% SDS, 0.2N NaOH) were added, well mixed and incubated for 7 min at room temperature. After the alkaline SDS treatment, 300 µl of ice-cold 3M sodium acetate pH 4.8 were added, and the mix is immediately spun at 13000g for 15 min at room temperature. The supernatant was transferred to a new 1.5 ml conical centrifuge tube and 650 µl of isopropanol (stored at room temperature) were added. The mix was then centrifuged at 13,000 x g for 5 min. The supernatant fluid was discarded, the pellet washed with 70% ethanol, and resuspended in 320 µl sterile distilled water.

The presence of individual phage 77 ORF DNA inserts in the plasmid was verified by PCR amplification using 1.5 µl transformant miniprep DNA in a PCR with primers flanking the cloning site of ORF in pTHA vector (HAF and HAR). The composition of the PCR reaction and the cycling parameters are identical to those employed for library screening described above.

Example 5. Functional assay for bacterial inhibitory activity of bacteriophage 77 ORFs.

The anti-microbial activity of individual phage 77 ORFs was monitored by two growth inhibitory assays, one on solid agar medium, the other in liquid medium.

In general, *Staphylococcus* bacteria transformed with expression plasmids containing individual ORFs were grown in normal TSA medium and stored in 19% glycerol. At pre-determined times, arsenite was added to the culture to induce transcription of the phage 77 ORFs cloned immediately downstream from an arsenite-inducible promoter in the pTHA expression plasmid.

The effect of ORF induction on bacterial growth characteristics was then monitored and quantitated. The growth inhibition assay on solid medium was performed by streaking pTHA/ORF containing *S. aureus* transformant onto LB-Kn and TSA-Kn plates containing increasing concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M). Arsenite is used to induce the expression of cloned DNA in pTHA vector. In parallel, 3 μ l of 1/10 and 1/100 dilutions of the frozen cultures of the pTHA/ORF transformants were spotted as single drops onto LB-Kn and TSA-Kn plates containing increasing concentration of sodium arsenite (0; 2.5; 5; and 7.5 μ M). The plates were then incubated 16 hrs at 37°C, and the effect of arsenite-induced ORF expression on bacterial growth was monitored and quantitated by comparing the extent to that seen in control plates. As positive controls for growth inhibition, the *holin/lysin* genes of the *Staphylococcus aureus* phage Twort (Loessner et al., 1998) was subcloned into the pTHA *ars* inducible vector and used.

For the growth inhibition assay in liquid medium, stationary phase cultures were prepared by inoculating 2.5ml TSB-Kn with frozen *S. aureus* RN4220 transformants containing phage 77 ORFs cloned in pTHA vector followed by incubation for 16 hrs at 37°C. These cultures were then diluted 1/100 in the same medium, and the bacteria were allowed to grow for 2 hrs at 37°C to reach early log phase. 150 μ l of such culture were then mixed with 2.35 ml TSB-Kn medium with or without arsenite (the final concentration of arsenite in the medium was 0 or 5 μ M arsenite). After 3.5 hrs incubation at 37°C with shaking at 250 rpm, 100 μ l of bacterial culture was removed from each tube for OD₅₆₅ measurement. Serial ten-fold dilutions of the culture in buffered saline solution (0.85% NaCl) were then spotted onto TSB-Kn plates. The plates were incubated at 37°C 16 hrs and the number of surviving colonies counted the following day. The growth inhibitory property of individual ORFs was then quantitated by comparing CFU numbers under normal or arsenite-induction conditions. A schematic flow of the inhibition analysis is shown in Fig. 3 (also applicable to inhibition analysis for the other phage and bacteria pointed out herein). Inhibition results are shown in Figures 4A-C.

Example 6: Identification of Cecropin Signature Motif in *Staphylococcus aureus* Bacteriophage 3A ORF

The genome for *S. aureus* bacteriophage 3A was determined and the sequence was analyzed essentially as described for bacteriophage 77 in the examples above. Upon blast analysis of the identified open reading frames of phage 3A, the presence of an amino acid sequence corresponding to a cecropin signature motif was observed.

- 5 This motif (WDGHKTLEK) is located at position aa 481-489. Cecropins were originally identified in proteins from the cecropia moth and are recognized as potent antibacterial proteins that constitute an important part of the cell-free immunity of insects. Cecropins are small proteins (31-39 amino acid residues) that are active against both Gram-positive and Gram-negative bacteria by disrupting the bacterial
10 membranes. Although the mechanisms by which the cecropins cause cell death are not fully understood, it is generally thought to involve channel formation and membrane destabilization.

- The identification of a motif corresponding to a known inhibitor suggests that the product of ORF002 is also an inhibitory compound. Such inhibitory activity can
15 be confirmed as described herein or by other methods known in the art. Confirmation of the inhibitory activity would indicate that the ORF product could serve as the basis for construction of mimetic compounds and other inhibitors directed to the target of the ORF002 product.

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- 20 Boman, 1991, *Cell* 65:205-207.

Boman et al., 1991, *Eur. J. Biochem.* 201:23-31.

Wang et al., *J. Biol. Chem.* 273:27438-27448.

Example 7. Growth of *Staphylococcus aureus* bacteriophage 44AHJD:

- 25 *Staphylococcus aureus* propagating strain (PS 44A) (Felix d'Herelle Reference Centre #HER 1101) was used as a host to propagate its respective phage 44AHJD (Felix d'Herelle Reference Centre #HER 101). Two rounds of plaque purification of phage 44AHJD were performed on soft agar essentially as described in Sambrook *et al.* (1989). Briefly, the *Staphylococcus aureus* PS strain was grown overnight at 37°C
30 in Nutrient Broth [NB: 3 g Bacto Beef Extract, 5 g Bactopeptone per liter, (Difco Laboratories # 0003-17-8), supplemented with 0.5% NaCl]. The culture was then diluted 20 fold in NB and incubated at 37°C until an OD₅₄₀ of 0.2. In order to obtain single plaques, phage 44AHJD was subjected to 10-fold serial dilutions using the phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin) and 10 µl
35 were used to infect 0.5 ml of the cell suspension in the presence of 400 µg/ml of

CaCl₂. After incubation of 15 min at room temperature, 2 ml of melted soft agar (NB supplemented with 0.6% of agar) were added to the mixture and poured onto the surface of 100 mm nutrient agar plates (3 g Bacto Beef extract, 5 g Bactopeptone, 0.5% NaCl and 15 g of Bacto agar per liter (Difco Laboratories # 0001-17-0). After
5 overnight incubation at 37°C, a single plaque was isolated, resuspended in 1ml of phage buffer by end over end rotation for 2 h at room temperature and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock.

Large scale purification of bacteriophage and preparation of phage DNA was
10 as follows.

The propagation method was carried out by using the agar layer method described by Swanstörn and Adams (1951). Briefly, the PS 44A strain was grown to stationary phase overnight at 37°C in Nutrient Broth. The culture was then diluted 20x in NB and incubated at 37°C until the A₅₄₀ = 0.2. The suspension (15x10⁷ Bacteria)
15 was then mixed with 15x10⁵ phage particles to give a ratio of 100-bacteria/phage particle in the presence of 400 µg/ml of CaCl₂. After incubation of 15 min at room temperature, 7.5 ml of melted soft agar were added to the mixture and poured onto the surface of 150 mm nutrient agar plates and incubated overnight at 37°C. To collect the lysate, 20 ml of NB were added to each plate and the soft agar layer was collected by
20 scrapping off with a clean microscope slide and shaken vigorously for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) is collected and subjected to a treatment with 10 µg/ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, 10% (w/v) of PEG 8000 and 0.5 M of NaCl were
25 added to the lysate and the mixture was incubated on ice for 16 h. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman).

The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1
30 volume of chloroform and further purified by centrifugation on a preformed cesium chloride step gradient as described in Sambrook *et al.* (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 h at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 x g) for 24 h at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 h at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 h at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

Example 8. DNA sequencing of the Bacteriophage 44 AHJD genome.

Four mg of phage DNA was diluted in 200 µl of TE pH 8.0 in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4 cycles and size fractionated on 1% agarose gels. The sonicated DNA was then size fractionated by gel electrophoresis. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen) and eluted in 50 µl of 1mM Tris-HCl [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of *E. coli* DNA polymerase 1 as follows. Reactions were performed in a final volume of 100 µl containing DNA, 10 mM Tris-HCl pH 8.0, 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 5 µg BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of Klenow fragment (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was ethanol precipitated and resuspended in 20 µl of H₂O.

Cloning of the sonicated phage DNA into pKSII vector and transformation:

Blunt-ended DNA fragments were cloned by ligation directly into the *HincII* site of the pKSII vector (Stratagene) dephosphorylated with calf intestinal alkaline phosphatase (New England Biolabs). A typical reaction contained 100 ng of vector, 2

to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) overnight at 16°C. Transformation and selection of positive clones was performed in the host strain DH10 β of *E. coli* using ampicillin as a selective antibiotic as described in Sambrook *et al.* (1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 µl LB and 100 µg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *HincII* cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 µl reaction volume containing 10 mM Tris-HCl (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 mM primer, 187.5 µM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism BigDye™ primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction kit.

Example 9. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI

prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Staphylococcus aureus* bacteriophage 44AHJD is shown in Table 16.

A software program was used on the assembled sequence of bacteriophage 44AHJD to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(<http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c>) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage 44AHJD are listed in Tables 17 & 18.

Sequence homology searches for each ORF were carried out using an implementation of blast programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- 25 iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) *Staphylococcus aureus* NCTC 8325 (<ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa>);
- vi) *Staphylococcus pyogenes* (ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- 30 vii) PRODOM (ftp://ftp.toulouse.inra.fr/pub/prodom/current_release/prodom99_1.forblast.gz);
- viii) DOMO (<ftp://ftp.infobiogen.fr/pub/db/domo/>);

ix) TREMBL (ftp://www.expasy.ch/databases/sp_tr_nrdb/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 44AHJD are shown in Tables 19 & 20.

5 Example 10. Sub-Cloning of Bacteriophage 44 AHJD ORFs.

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 44 AHJD ORF sequence is inducible. For example, the shuttle vector pT0021, in which the firefly luciferase (*lucFF*) expression is controlled by the *ars* (arsenite) promoter/operator (Tauriainen et al., 1997), can be modified in the following fashion. Two oligonucleotides corresponding to a short antigenic peptide derived from the hemagglutinin protein of influenza virus (HA epitope tag) were synthesized (Field et al., 1988). The sense strand HA tag sequence (with *Bam*HI, *Sal*I and *Hind*III cloning sites) is:

5'-gatcccggtcgaccaagcttTACCCATACGACGTCCCAGACTACGCCAGCTGA-3'

15 (where upper case letters denote the nucleotide sequence of the HA tag); the antisense strand HA tag sequence (with a *Hind*III cloning site) is:

5'-agctTCAGCTGGCGTAGTCTGGGACGTCGTATGGGTAAagcttggtcgaccgg-3'

(where upper case letters denote the sequence of the HA tag). The two HA tag oligonucleotides were annealed and ligated into pT0021 vector which had been digested with *Bam*HI and *Hind*III. This manipulation resulted in replacement of the *lucFF* gene by the HA tag. This modified shuttle vector containing the *arsenite* inducible promoter, the *arsR* gene, and HA tag was named pTHA. A diagram outlining our modification of pT0021 to generate pTHA is shown in Fig. 1A (another useful vector construct is shown in Fig. 1B).

25 Each ORF, encoded by Bacteriophage 44 AHJD, larger than 33 amino acids and having a Shine-Dalgarno sequence upstream of the initiation codon can be selected for functional analysis for bacterial inhibition. Each individual ORF, from initiation codon to last codon (excluding the stop codon), can be amplified from phage genomic DNA using the polymerase chain reaction (PCR). For PCR amplification of ORFs, each sense strand primer targets the initiation codon and is preceded by a *Bam*HI restriction site (5'-cgggatcc-3') and each antisense oligonucleotide targets the pentultimate codon (the one before the stop codon) of the ORF and is preceded by a *Sal*I restriction site (5'-gcgtcgaccg-3'). The PCR product of each ORF can be gel

purified and digested with *Bam*HI and *Sa*II. The digested PCR product can then be gel purified using the Qiagen kit as described, ligated into *Bam*HI and *Sa*II digested pTHA vector, and used to transform *E. coli* bacterial strain DH10 β (as described above). As a result of this manipulation, the HA tag is set inframe with the ORF and is positioned at the carboxy terminus of each ORF (pTHA/ORF clones). Recombinant pTHA/ORF clones will be picked and their insert sizes were confirmed by PCR analysis using primers flanking the cloning site. The following primers can be used for PCR amplification: HAF: 5'TATTATCCAAAACCTTGAACA^{3'}; HAR: 5'CGGTGGTATATCCAGTGATT^{3'}. The sequence integrity of cloned ORFs can be verified directly by DNA sequencing using primers HAF and HAR. In cases where verification of ORF sequence can not be achieved by one pass with the sequencing primers, additional internal primers will be selected and used for sequencing.

Staphylococcus aureus strain RN4220 (Kreiswirth et al., 1983) will be used as a recipient for the expression of recombinant plasmids. Electroporation will be performed essentially as previously described (Schenk and Laddaga, 1992). Selection of recombinant clones will be performed on Luria-Broth agar (LB-agar) plates containing 30 μ g/ml of kanamycin.

Alternatively, a constitutive promoter can be used to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids will be introduced into *Staphylococcus aureus* strain RN4220 (Kreiswirth et al., 1983) using electroporation as previously described (Schenk and Laddaga, 1992).

Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), can be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10. Recombinant clones are then picked and their insert sizes confirmed by

PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs can be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal
5 primers can be selected and used for sequencing. Recombinant plasmids can be introduced into *Staphylococcus aureus* strain RN4220 (Kreiwirth et al., 1983) using electoporation as previously described (Schenk and Laddaga, 1992).

Induction of gene expression from the *ars* promoter.

If an inducible promoter is used, e.g., the *ars* promoter, induction can be
10 assessed, for example, in either of the two methods.

1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of *S. aureus* transformed cells containing phage 44 AHJD ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M). The
15 plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are
20 then diluted to the mid log phase ($OD_{540}=0.2$) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μ l/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μ M) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage 44 AHJD ORFs on bacterial cell growth is then monitored by measuring the OD_{540} and comparing the
25 rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the *Staphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) can be
30 subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of

colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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Example 11. Growth of *Enterococcus* bacteriophage 182 and purification of genomic DNA.

The *Enterococcus* propagating strain (PS) (*Enterococcus* sp. Group D, Felix d'Herelle Reference Centre #HER 1080) was used as host to propagate its respective
10 phage 182 (Felix d'Herelle Reference Centre #HER 80). Two rounds of plaque purification of phage 182 were performed on soft agar essentially as described in Sambrook *et al.* (1989). Briefly, the *Enterococcus* sp. PS strain was grown overnight at 37°C in Tryptic Soy Broth [TSB: 17 g Bacto tryptone, 3 g Bacto soytone, 2.5 g Bacto dextrose, 5 g Sodium chloride, and 2.5 g Dipotassium phosphate per liter
15 (Difco Laboratories (#0370-17-3))]. The culture was then diluted 20 fold in TSB and incubated at 37°C until the $OD_{540} = 0.2$ (early log phase) with constant agitation. In order to obtain single plaques, phage 182 was subjected to 10 fold serial dilutions using the phage buffer (1 mM $MgSO_4$, 5 mM $MgCl_2$, 80 mM NaCl and 0.1% Gelatin (w/v)) and 10 l of each dilution was used to infect 0.5 ml of the bacterial cell
20 suspension. After incubation at 15 min at 37°C, 2 ml of melted soft agar (TSB supplemented with 0.6% agar) was added to the mixture and poured onto the surface of 100 mm Tryptic Soy Agar plates [TSA: 15 g Tryptone peptone, 5 g Soytone peptone, 5 g Sodium chloride and 15 g of Agar per liter (Difco Laboratories #0369-17)]. After overnight incubation at 37°C, a single plaque was isolated, resuspended in
25 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

The propagation procedure for bacteriophage 182 was modified from the agar
30 layer method of Swanstörn and Adams (1951). Briefly, the *Enterococcus* sp. PS strain was grown to stationary phase overnight at 37°C in TSB. The culture was then diluted 20 fold in TSB and incubated at 37°C until the $A_{540} = 0.2$. The suspension (15×10^7 Bacteria) was then mixed with 15×10^5 plaque forming units (pfu) to give a

ratio of 100-bacteria/pfu. After incubation of 15 min at 37°C, 7.5 ml of melted soft agar (TSB plus 0.6% agar) were added to the mixture and poured onto the surface of 150 mm TSA plates and incubated 16 hrs at 37°C. To collect the plate lysate, 20 ml of TSB were added to each plate and the soft agar layer was collected by scrapping off with a clean microscope slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant fluid (lysate) is collected and subjected to a treatment with 10 µg /ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (1 mM MgSO₄, 5 mM MgCl₂, 80 mM NaCl and 0.1% Gelatin). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a cesium chloride step gradient as described in Sambrook *et al.* (1989), using a TLS 55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phages were harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 g/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

Example 12. DNA sequencing of the Bacteriophage 182 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 s spaced by 15 s cooling in ice/water for 3 to 4

cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of *E. coli* DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of the Klenow large fragment of DNA polymerase I (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in 20 µl of H₂O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 µl LB and 100 µg/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 µl reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 µM primer, 187.5 µM each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec

denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

5 The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and
10 the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction kit.

15

Example 13. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI
20 prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Enterococcus* bacteriophage 182 is shown in Table 21.

A software program was used on the assembled sequence of bacteriophage 182 to identify all putative ORFs larger than 33 codons. The software scans the primary
25 nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG, GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI([http://www.ncbi.nlm.nih.gov/htbin-](http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c)
30 [post/Taxonomy/wprintgc?mode=c](http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c)) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the

next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is

5 performed on all three reading frames of both DNA strands of the phage sequence.

The predicted ORFs for bacteriophage 182 are listed in Tables 22 & 23.

Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:

- 10 (i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) staphylococcus aureus NCTC 8325 ([ftp://ftp.genome.ou.edu/pub/staph/staph-](ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa)
- 15 [1k.fa](ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa));
- vi) streptococcus pyrogenes
(ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- vii) PRODOM
(ftp://ftp.toulouse.inra.fr/pub/prodom/current_release/prodom99.1.forblast.gz);
- 20 viii) DOMO (<ftp://ftp.infobiogen.fr/pub/db/domo/>);
- ix) TREMBL (ftp://www.expasy.ch/databases/sp_tr_nrd/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage 182 are shown in Tables 24 & 26.

25 Example 14. Sub-Cloning of Bacteriophage 182 ORFs.

Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage 182 ORF sequence is inducible.

For example, the plasmid pND50 replicates in *E. coli*, *E. faecalis*, and *S. aureus*

- 30 (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. *Antimicrob. Agents Chemother.* 40, 1157-1163). This plasmid can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the firefly luciferase (*lucFF*)

expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997). Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain the *ars* promoter, *arsR* gene
5 and a cloning site for introduction of individual phage ORFs downstream from a shine-dalgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system. The *nisA* promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal
10 transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the *nisA* promoter (10- to 60-fold induction) can be obtained in all of the
15 species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transcription in *Enterococcus*.

Alternatively, a constitutive promoter can be used (e.g., the β -lactamase
20 promoter is constitutive in *E. faecalis* – see ref. 1) to drive expression of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *E. faecalis* strain FA2-2 by electroporation, as previously described (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S.,
25 and Inoue, M. 1996. *Antimicrob. Agents Chemother.* 40, 1157-1163).

Cloning of ORFs with a Shine-Dalgarno sequence

ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop
30 codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on

the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10 β . Recombinant clones are then picked and their insert sizes confirmed by PCR analysis using primers flanking the cloning site as well as restriction digestion.

- 5 The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into *E. faecalis* strain FA2-2 by electroporation, as previously described
- 10 (Yamagishi, J., Kojima, T., Oyamada, Y., Fujimoto, K., Hattori, H., Nakamura, S., and Inoue, M. 1996. *Antimicrob. Agents Chemother.* 40, 1157-1163).

Induction of gene expression from the *ars* promoter.

If an inducible promoter is used, e.g., the *ars* promoter, induction can be assessed, for example, in either of the two methods.

15 1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of *E. faecalis* transformed cells containing phage 182 ORF onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M). The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF

- 20 transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

- Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase ($OD_{540}=2$) with fresh media containing antibiotic
- 25 and transferred to 96-well microtitration plates (100 μ l/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μ M) and the culture incubated for an additional 2 h at 37°C. The effect of expression of the phage 182 ORFs on bacterial cell growth is then monitored by measuring the OD_{540} and comparing the rate of growth to the culture not containing inducer. As positive controls for growth
- 30 inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the *Staphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G.,

Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) were subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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Example 15. Growth of *Streptococcus* bacteriophage Dp-1 and purification of genomic DNA.

The *Streptococcus pneumoniae* R6 propagating strain (PS) (Tomasz, 1966) was used as host to propagate its respective phage Dp-1 (McDonnell et al., 1975). (Alternatively, *Streptococcus (Diplococcus) pneumoniae* R36A could be used. Strain R36A is available from ATCC as #11733 or 27336. *Streptococcus pneumoniae* is also available from Felix d'Herelle Reference Center in Quebec, Canada as catalog number HER 1054. Other *S. pneumoniae* strains are also available from ATCC.) Two rounds of plaque purification of phage Dp-1 were performed on soft agar essentially as described in Sambrook et al. (1989). Briefly, the *Streptococcus* R6 PS strain was grown overnight at 37°C in K-Cat media [K-Cat: 10 g Bacto casitone, 5 g Bacto tryptone, 1 g Yeast extract, 5g Potassium chloride, 0.2% Glucose, 30mM Potassium phosphate buffer [pH 8] and 250,000 Units Catalase per liter (Boehringer Mannheim #10683600). The culture was then diluted 20 fold in K-CAT and

incubated at 37°C until the $OD_{540} = 0.2$ (early log phase) with constant agitation. In order to obtain single plaques, Dp-1 phage was subjected to 10-fold serial dilutions using the phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM $MgCl_2$) and 10 μ l of each dilution was used to infect 0.5 ml of the cell suspension.

5 After incubation of 15 min at 37°C, 2 ml of melted soft agar (K-CAT supplemented with 0.8% of agar) were added to the mixture and poured onto the surface of 100 mm K-CAT agar plates [K-CAT supplemented with 1.2 % of agar]. After solidification of the soft agar layer, an additional 5 ml of melted soft agar was added to visualize distinct plaques (Ronda et al., 1978). After overnight incubation at 37°C, a single

10 plaque was isolated, resuspended in 1 ml of phage buffer by end over end rotation for 2 hrs at room temperature, and the phage suspension was diluted and used for a second infection as described above. After overnight incubation at 37°C, a single plaque was isolated and used as a stock for all subsequent manipulations.

The propagation procedure for bacteriophage Dp-1 was modified from the

15 agar layer method of Swanstörn and Adams (1951). Briefly, the R6 strain of *Streptococcus pneumoniae* was grown to stationary phase overnight at 37°C in K-CAT. The culture was then diluted 20 fold in K-CAT and incubated at 37°C until the $OD_{540} = 0.2$. The suspension (15×10^7 Bacteria) was then mixed with 15×10^5 plaque forming units (pfu) to give a ratio of 100-bacteria/pfu. After incubation of 15 min at

20 37°C, 7.5 ml of melted soft agar (K-CAT plus 0.8% agar) were added to the mixture and poured onto the surface of 150 mm K-CAT agar plates and incubated 16 hrs at 37°C. After solidification of the soft agar layer, 7.5 ml of melted soft agar were added to each plate. To collect the plate lysate, 20 ml of K-CAT media were added to each plate and the soft agar layers were collected by scrapping off with a clean microscope

25 slide followed by vigorous shaking of the agar suspension for 5 min to break up the agar. The mixture was then centrifuged for 10 min at 4,000 rpm (2,830 xg) using a JA-10 rotor (Beckman) and the supernatant (lysate) was collected and subjected to a treatment with 10 μ g /ml of DNase I and RNase A for 30 min at 37°C. To precipitate the phage particles, the phage suspension was adjusted to 10% (w/v) of PEG 8000 and

30 0.5 M of NaCl followed by incubation at 4°C for 16 hrs. The phage was recovered by centrifugation at 4,000 rpm (3,500 xg) for 20 min at 4°C on a GS-6R table top centrifuge (Beckman). The pellet was resuspended with 2 ml of phage buffer (100 mM Tris-HCl [pH 7.5], 100 mM NaCl and 10 mM $MgCl_2$). The phage suspension was extracted with 1 volume of chloroform and further purified by centrifugation on a

35 cesium chloride step gradient as described in Sambrook et al. (1989), using a TLS-55 rotor and centrifuged in an Optima TLX ultracentrifuge (Beckman) for 2 hrs at 28,000 rpm (67,000 xg) at 4°C. Banded phage was collected and ultracentrifuged again on an

isopycnic cesium chloride gradient (1.45 g/ml) at 40,000 rpm (64,000 xg) for 24 hrs at 4°C using a TLV rotor (Beckman). The phage was harvested and dialyzed for 4 hrs at room temperature against 4 L of dialysis buffer consisting of 10 mM NaCl, 50 mM Tris-HCl [pH 8] and 10 mM MgCl₂. Phage DNA was prepared from the phage suspension by adding 20 mM EDTA, 50 µg/ml Proteinase K and 0.5% SDS and incubating for 1 hr at 65°C, followed by successive extractions with 1 volume of phenol, 1 volume of phenol-chloroform and 1 volume of chloroform. The DNA was then dialyzed overnight at 4°C against 4 L of TE (10 mM Tris-HCl [pH 8.0], 1mM EDTA).

Example 16. DNA sequencing of the Bacteriophage Dp-1 genome.

Four micrograms of phage DNA was diluted in 200 µl of TE (10 mM Tris, [pH 8.0], 1 mM EDTA) in a 1.5 ml eppendorf tube and sonication was performed (550 Sonic Dismembrator, Fisher Scientific). Samples were sonicated under an amplitude of 3 µm with bursts of 5 sec spaced by 15 sec cooling in ice/water for 3 to 4 cycles. The sonicated DNA was then size fractionated by electrophoresis on 1% agarose gels utilizing TAE (1 x TAE is: 40 mM Tris-acetate, 1 mM EDTA [pH 8.0]) as the running buffer. Fractions ranging from 1 to 2 kbp were excised from the agarose gel and purified using a commercial DNA extraction system according to the instructions of the manufacturer (Qiagen), with a final elution of 50 µl of 1 mM Tris [pH 8.5].

The ends of the sonicated DNA fragments were repaired with a combination of T4 DNA polymerase and the Klenow fragment of *E. coli* DNA polymerase I, as follows. Reactions were performed in a reaction mixture (final volume, 100 µl) containing sonicated phage DNA, 10 mM Tris-HCl [pH 8.0], 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA, 100 µM of each dNTP and 15 units of T4 DNA polymerase (New England Biolabs) for 20 min at 12°C followed by addition of 12.5 units of the Klenow large fragment of DNA polymerase I (New England Biolabs) for 15 min at room temperature. The reaction was stopped by two phenol/chloroform extractions and the DNA was precipitated with ethanol and the final DNA pellet resuspended in 20 µl of H₂O.

Blunt-ended DNA fragments were cloned by ligation directly into the *Hinc* II site of the pKSII+ vector (New England Biolabs) dephosphorylated by treatment with calf intestinal alkaline phosphatase (New England Biolabs). A typical ligation reaction contained 100 ng of vector DNA, 2 to 5 µl of repaired sonicated phage DNA (50-100 ng) in a final volume of 20 µl containing 800 units of T4 DNA ligase (New England Biolabs) and was incubated overnight at 16°C. Transformation and selection

of bacterial clones containing recombinant plasmids was performed in *E. coli* DH10 β according to standard procedures (Sambrook *et al.*, 1989).

Recombinant clones were picked from agar plates into 96-well plates containing 100 μ l LB and 100 μ g/ml ampicillin and incubated at 37°C. The presence of phage DNA insert was confirmed by PCR amplification using T3 and T7 primers flanking the *Hinc* II cloning site of the pKS vector. PCR amplification of the potential foreign inserts was performed in a 15 μ l reaction volume containing 10 mM Tris (pH 8.3), 50 mM KCl, 1.5 mM MgCl₂, 0.02% gelatin, 1 μ M primer, 187.5 μ M each dNTP, and 0.75 units *Taq* polymerase (BRL). The thermocycling parameters were as follows: 2 min initial denaturation at 94°C for 2 min, followed by 20 cycles of 30 sec denaturation at 94°C, 30 sec annealing at 58°C, and 2 min extension at 72°C, followed by a single extension step at 72°C for 10 min. Clones with insert sizes of 1 to 2 kbp were selected and plasmid DNA was prepared from the selected clones using the QIAprep™ spin miniprep kit (Qiagen).

The nucleotide sequence of the extremities of each recombinant clone was determined using an ABI 377-36 automated sequencer with two types of chemistry: ABI prism Big Dye™ primer cycle sequencing (21M13 primer: #403055)(M13REV primer: #403056) or ABI prism Big Dye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). To ensure co-linearity of the sequence data and the genome, all regions of the phage genome were sequenced at least once from both directions on two separate clones. In areas that this criteria was not initially met, a sequencing primer was selected and phage DNA was used directly as sequencing template employing ABI prism Big Dye™ terminator cycle sequencing ready reaction kit.

Example 17. Bioinformatic management of primary nucleotide sequence.

Sequence contigs were assembled using Sequencher™ 3.1 software (GeneCodes). To close contig gaps, sequencing primers were selected near the edge of the contigs. Phage DNA was used directly as sequencing template employing ABI prism BigDye™ terminator cycle sequencing ready reaction kit (Applied Biosystems; #4303152). The complete sequence of *Streptococcus* bacteriophage Dp-1 is shown in Table 28.

A software program was used on the assembled sequence of bacteriophage Dp-1 to identify all putative ORFs larger than 33 codons. The software scans the primary nucleotide sequence starting at nucleotide #1 for an appropriate start codon. Three possible selections can be made for defining the nature of the start codon; I) selection of ATG, II) selection of ATG or GTG, and III) selection of either ATG,

GTG, TTG, CTG, ATT, ATC, and ATA. This latter initiation codon set corresponds to the one reported by the NCBI(<http://www.ncbi.nlm.nih.gov/htbin-post/Taxonomy/wprintgc?mode=c>) for the bacterial genetic code. When an appropriate start codon is encountered, a counting mechanism is employed to count the number of codons (groups of three nucleotides) between this start codon and the next stop codon downstream of it. If a threshold value of 33 is reached, or exceeded, then the sequence encompassed by these two codons is defined as an ORF. This procedure is repeated, each time starting at the next nucleotide following the previous stop codon found, in order to identify all the other putative ORFs. The scan is performed on all three reading frames of both DNA strands of the phage sequence. The predicted ORFs for bacteriophage Dp-1 are listed in Tables 29 and 30, and Fig. 6.

Sequence homology searches for each ORF were carried out using an implementation of BLAST programs. Downloaded public databases used for sequence analysis include:

- (i) non-redundant GenBank (<ftp://ncbi.nlm.nih.gov/blast/db/nr.Z>),
- ii) Swissprot (<ftp://ncbi.nlm.nih.gov/blast/db/swissprot.Z>);
- iii) vector (<ftp://ncbi.nlm.nih.gov/blast/db/vector.Z>);
- iv) pdbaa databases (<ftp://ncbi.nlm.nih.gov/blast/db/pdbaa.Z>);
- v) staphylococcus aureus NCTC 8325
(<ftp://ftp.genome.ou.edu/pub/staph/staph-1k.fa>);
- vi) streptococcus pyogenes
(ftp://ftp.tigr.org/pub/data/s_pneumoniae/gsp.contigs.112197.Z);
- vii) PRODOM
(ftp://ftp.toulouse.inra.fr/pub/prodom/current_release/prodom99.1.forblast.gz);
- viii) DOMO (<ftp://ftp.infobiogen.fr/pub/db/domo/>);
- ix) TREMBL (ftp://www.expasy.ch/databases/sp_tr_nrd/fasta/)

The results of the homology searches performed on the ORFs of bacteriophage Dp-1 are shown in Table 31.

Example 18. Sub-Cloning of Bacteriophage Dp-1 ORFs.

Preparation of the shuttle expression vector

Expression preferably utilizes a shuttle expression vector which is arranged such that expression of the exogenous bacteriophage Dp-1 ORF sequence is inducible. For example, the plasmid pLSE4 replicates in *E. coli*, and *S. pneumoniae* (Diaz and Garcia, 1990). This plasmid can be modified by conventional techniques to insert the inducible arsenite promoter, derived from the shuttle vector pT0021, in which the

firefly luciferase (*lucFF*) expression is controlled by the *ars* promoter/operator from a *S. aureus* plasmid (Tauriainen, S., Karp, M., Chang, W and Virta, M. (1997).

Recombinant luminescent bacteria for measuring bioavailable arsenite and antimonite. *Appl. Environ. Microbiol.* 63:4456-4461). This modified shuttle vector will contain

- 5 the *ars* promoter, *arsR* gene and a cloning site for introduction of individual phage ORFs downstream from a shine-dalgarno sequence.

Other inducible regulatory sequences can be utilized instead of the arsenite-inducible system. An example is a nisin-inducible system The *nisA* promoter activity is dependent on the proteins NisR and NisK, which constitute a two-component signal
10 transduction system that responds to the extracellular inducer nisin. The nisin sensitivity and inducer concentration required for maximal induction varies among the strains, but is functional in *Streptococcus pyogenes*, *Streptococcus agalactiae*, *Streptococcus pneumoniae*, *Enterococcus faecalis*, and *Bacillus subtilis*. Significant induction of the *nisA* promoter (10- to 60-fold induction) can be obtained in all of the
15 species. A vector containing this promoter was published as Eichenbaum Z, Federle MJ, Marra D, de Vos WM, Kuipers OP, Kleerebezem M, and Scott JR (1998) *Appl Environ Microbiol* 64, 2763-2769. Other vectors, e.g., plasmids, can also be utilized which will allow replication and transcription in *Streptococcus*.

Alternatively, a constitutive promoter can be used to drive expression
20 of the introduced ORF, and compare cell growth to control bacterial cells containing the parental vector lacking any introduced phage ORF. Recombinant plasmids are introduced into *S. pneumoniae* R6 as previously described (Diaz and Garcia, 1990)

Cloning of ORFs with a Shine-Dalgarno sequence

- 25 ORFs with a Shine-Dalgarno sequence are selected for functional analysis of bacterial killing. Each ORF, from initiation codon to last codon (excluding the stop codon), will be amplified by PCR from phage genomic DNA. For PCR amplification of ORFs, each sense strand primer starts at the initiation codon and is preceded by a restriction site and each antisense strand starts at the last codon (excluding the stop
30 codon) and is preceded by a different restriction site. The PCR product of each ORF will be gel purified and digested with the restriction enzymes with sites contained on the PCR oligonucleotides. The digested PCR product is then gel purified using the Qiagen kit, ligated into the modified shuttle vector, and used to transform bacterial strain DH10 β . Recombinant clones are then picked and their insert sizes confirmed
35 by PCR analysis using primers flanking the cloning site as well as restriction digestion. The sequence fidelity of cloned ORFs will be verified by DNA sequencing using the same primers as used for PCR. In the cases that the verification of ORFs can not be achieved by one path of sequencing using primers flanking the cloning site

internal primers will be selected and used for sequencing. Recombinant plasmids will be introduced into *S. pneumoniae* R6 as previously described (Diaz and Garcia, 1990).

Induction of gene expression from the *ars* promoter.

If an inducible promoter is used, e.g., the *ars* promoter, induction can be assessed, for example, in either of the two methods.

1. Screening on agar plates

The functional identification of killer ORFs can be performed by spreading an aliquot of *S. pneumoniae* transformed cells containing phage Dp-1 ORFs onto agar plates containing different concentrations of sodium arsenite (0; 2.5; 5; and 7.5 μ M).

The plates are incubated overnight at 37°C, after which a growth inhibition of the ORF transformants on plates that contain arsenite are compared to plates without arsenite.

2. Quantification of growth inhibition in liquid medium

Cells containing different recombinant plasmids can be grown for overnight at 37°C in LB medium supplemented with the appropriate antibiotic selection. These are then diluted to the mid log phase ($OD_{540}=2$) with fresh media containing antibiotic and transferred to 96-well microtitration plates (100 μ l/well). Inducer is then added at different final concentrations (ranging from 2.5 to 10 μ M) and the culture incubated for an additional 2 hrs at 37°C. The effect of expression of the phage Dp-1 ORFs on bacterial cell growth is then monitored by measuring the OD_{540} and comparing the rate of growth to the culture not containing inducer. [As positive controls for growth inhibition, the *kilA* gene of phage lambda (Reisinger, GR., Rietsch, A., Lubitz, W. and Blasi, U. 1993 *Virology* #193: 1033-1036), and the *holin/lysin* genes of the *Staphylococcus aureus* phage Twort (Loessner, MJ., Gaeng, S., Wendlinger, G., Maier, SK. and Scherer, S. 1998. *FEMS Microbiology Letters* #162:265-274) can be subcloned into the *ars* inducible vector. An aliquot of the induced and uninduced culture can also be plated out on agar plates containing an appropriate antibiotic selection but lacking inducer. Following incubation overnight at 37°C, the number of colonies is counted. Any ORF showing bacteriostatic activity will show a lower, but detectable, number of colonies on the agar plates when grown in the presence of inducer as compared to when grown in the absence of inducer. Any ORF showing full bacteriocidal activity will show no colonies on the agar plates, when grown in the presence of inducer as compared to when grown in the absence of inducer.

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10 All patents and publications mentioned in the specification are indicative of the levels of skill of those skilled in the art to which the invention pertains. All references cited in this disclosure are incorporated by reference to the same extent as if each reference had been incorporated by reference in its entirety individually.

15 One skilled in the art would readily appreciate that the present invention is well adapted to carry out the objects and obtain the ends and advantages mentioned, as well as those inherent therein. The specific methods and compositions described herein as presently representative of preferred embodiments are exemplary and are not intended as limitations on the scope of the invention. Changes therein and other uses will occur to those skilled in the art which are encompassed within the spirit of the invention are defined by the scope of the claims.

20 It will be readily apparent to one skilled in the art that varying substitutions and modifications may be made to the invention disclosed herein without departing from the scope and spirit of the invention. For example, those skilled in the art will recognize that the invention may suitably be practiced using a variety of different bacteria, bacteriophage, and sequencing methods within the general descriptions
25 provided.

30 The invention illustratively described herein suitably may be practiced in the absence of any element or elements, limitation or limitations which is not specifically disclosed herein. Thus, for example, in each instance herein any of the terms "comprising," "consisting essentially of" and "consisting of" may be replaced with either of the other two terms. The terms and expressions which have been employed are used as terms of description and not of limitation, and there is not intention that in the use of such terms and expressions of excluding any equivalents of the features shown and described or portions thereof, but it is recognized that various
35 modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the concepts herein disclosed may be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of this invention as defined by the appended claims.

In addition, where features or aspects of the invention are described in terms of Markush groups or other grouping of alternatives, those skilled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group or other group. For example, 5 if there are alternatives A, B, and C, all of the following possibilities are included: A separately, B separately, C separately, A and B, A and C, B and C, and A and B and C. Thus, for example, for the bacteria and phage specified herein, the embodiments expressly include any subset or subgroup of those bacteria and/or phage. While each such subset or subgroup could be listed separately, for the sake of brevity, such a 10 listing is replaced by the present description.

Thus, additional embodiments are within the scope of the invention and within the following claims.

Table 1

Phages against human and animal pathogenic bacteria

5

I. Pathogen name	Phage name	II. Catalog#	Origin/reference
<i>Acinetobacter calcoaceticus</i>	A3/2 A10/45 A36 B9GP B ₉ PP BS46 E13 E14 531		Felix d'Herelle Reference Centre, Quebec, Quebec
	Ap3 P78		J. Bacteriol 1984. 157: 179-183 J. Gen. Microbiol 1986.132: 2633-2636
<i>Acinetobacter haemolyticus</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Acinetobacter johnsonii</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Acinetobacter sp.</i>	BP1		J. Virol. 1968.2:716-722
	G4, HP2, HP3 & HP4		Can. J. Microbiol. 1966.12:1023-1030 & J. Virol. 1974.13:46-52 & Arch. Virol. 1994.135:345-354
	A1, A4, A9 & 196		Arch. Virol. 1994.135:345-354
	HP1		Can. J. Microbiol. 1966.12:1023-1030
	A19, A23, A29, A31, A33, A34, A3759 & 2845		J. Microsc (Paris) 1973.16:215-224 & CR. Hebdo Seances Acad. Sci. Ser D. Sci Natur (Paris) 278:1907-1909 & Arch. Virol. 1994.135:345-354 & Rev. Can. Biol. 1970.29:317-320
<i>Actinobacillus actinomycetecomitans</i>			FEMS Microbiol Lett 1994. 119:329-337

			Infec. Immun. 1982. 35: 343-349
			Mol.Gen.Genet 1998.258: 323-325
	Aap247		Oral Micriol. Immunol 1997.12: 40-46
<i>Actinomyces viscosus</i>		43146-B1	The American Type Culture Collection
			Infect.Immun.1985.48:228-233
			Infect.Immun.1988.56:54-59
			Plasmid 1997.37:141-153
<i>Aeromonas hydrophila</i>	PM2** & PM3		FEMS Microbiol.Lett. 1990.57:277-282
	Aeh1 Aeh2 PM4 PM5 PM6 T7-ah		Felix d'Herelle Reference Centre, Quebec, Quebec

<i>Aeromonas salmonicida</i>	3 25 29 31 32 40RR _{2,8t} 43 51 56 59.1 65 Asp37		Felix d'Herelle Reference Centre, Quebec, Quebec
	55R.1		Can. J. Microbiol. 1983. 29: 1458-1461
<i>Alteromonas espejiana</i>	PM2**	27025-B1	The American Type Culture Collection
<i>Asticcacaulis biprosthecum</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Asticcacaulis excentricus</i>		15261-B1 15261-B2 15261-B3	The American Type Culture Collection
	φAc21 φAc24		
<i>Azotobacter vinelandii</i>	A14 A21 A31 A41 PAV1	12518-B1 12518-B4 12518-B5 12518-B9 12518-B10 13705-B1	The American Type Culture Collection
<i>Azotobacter sp.</i>			Virology 1972.49:439-452
<i>Bacteroides fragilis</i>	Bf-1		Rev. Infect. Dis. 1979. 1: 325-336
	B40-8		FEMS Microbiol. Lett. 1991. 66: 61-67
	HSP40		Appl. Environ. Microbiol. 1989. 55: 2696-2701
	phiA1		Zentralbl.bakteriol.1972.222:57-63
<i>Bdellovibrio bacteriovorus</i>	MAC-1		J. Gen. Microbiol. 1987. 133: 3065-3070
<i>Bdellovibrio sp.</i>	VL-1		J.Virol.1973.12:1522-1533
<i>Bordetella brochiseptica</i>	214		Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-13

<i>Bordetella parapertussis</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
			Mol. Gen. Mikrobiol. Virusol. 1988.4: 22-25
			Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-13
	41405		Zh.Mikrobiol.Epidemiol.Immuno. 1987.5:9-13
<i>Brucella abortus</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
	10/I 24/II 212/XV	23448-B1 23448-B2 23448-B3 17385-B1 17385-B2	The American Type Culture Collection
	BK-2, TB & Fj ^{oa}		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2: 48-52
	R/c & R/O		Dev. Biol. Stand. 1984.56: 55-62
<i>Brucella canis</i>	R/c		Dev. Biol. Stand. 1984.56: 55-62
<i>Brucella melitensis</i>	BK-2	23456-B1	The American Type Culture Collection
<i>Brucella suis</i>	Wb		Zentralbl.Veterinarmed.1975.22:866-867

	Fi** & TB		Zh.Mikrobiol.Epidemiol.Immunobiol.1983.2: 48-52
<i>Brucella sp.</i>			Can. J. Vet. Res. 1989.53: 319-325
			Res. Vet. Sci. 1988. 44: 45-49
	R		Zh.Mikrtobiol.Epidemiol.Immunobiol.1983.2: 48
<i>Campylobacter coli</i>		43133-B1	The American Type Culture Collection
<i>Campylobacter coli</i> (Cont'd)	18	43134-B1	The American Type Culture Collection
19		43135-B1	
20		43136-B1	
<i>Campylobacter jejuni</i>	1	35918-B1	The American Type Culture Collection
	2	35919-B1	
	3	35920-B1	
	4	35921-B1	
	5	35918-B2	
	6	35920-B2	
	7	35922-B2	
	8	35923-B1	
	9	35924-B1	
	10	35925-B1	
	11	35925-B2	
	12	35922-B2	
	13	35924-B2	
	14	35922-B3	
	17	43133-B1	
	18	43134-B1	
	19	43135-B1	
	20	43136-B1	
<i>Campylobacter</i> (<i>Helicobacter</i>) <i>pylori</i>	HP1		J. Med. Microbiol.1993. 38: 245-249
<i>Chlamydia psittaci</i>	Chp1**		J. Gen. Virol. 1989. 70: 3381-3390
<i>Clostridium</i> <i>acetobutylicum</i>	CAK-1		J.Bacteriol.1993.175:3838-3843

<i>Clostridium botulinum</i>			Nucleic Acids Res.1990.18:1291
			Bioch.Biophys.res.Commun.1990.171.1304-1311
			Microbiol.immunol.1981.25:915-927
			J.Vet.Med.Sci.1992.54:675-684
	CE β & CE γ		
<i>Clostridium difficile</i>	41 & 56		J. Clini.Microbiol. 1985.21:251-254

<i>Clostridium perfringens</i>			Rev.Can.Biol.1977.36:205-215
			FEMS Microbiol.Lett. 1990.54:323-326
<i>Clostridium sporogenes</i>	59 70 71 72S 72L	8074-B1 17886-B1 17886-B3 17886-B4 17886-B5 17886-B6	The American Type Culture Collection
<i>Clostridium tetani</i>	A & B		Rev.Can.Biol.1978.37:43-46
<i>Corynebacterium diphtheriae</i>			Vopr.Virusol.1986.31:577-584
<i>Corynebacterium pseudotuberculosis</i>	NN	12319-B1	The American Type Culture Collection
<i>Corynebacterium sp</i>	DLC 2921/49	12052-B1	The American Type Culture Collection

<i>Enterococcus faecalis</i>	42	19948-B1	The American Type Culture Collection
<i>Enterococcus faecium</i>	124 133	19950-B1 19953-b2 19953-B1	The American Type Culture Collection

<i>Escherichia coli</i>		11303-B14	The American Type Culture Collection
		11303-B10	
		11303-B21	
		8677-B1	
		11303-B13	
		13706-B4	
<i>Escherichia coli</i> (Cont'd)		15766-B1	The American Type Culture Collection
		15766-B1	
		1242-B5	
		15669-B2	
		15767-B1	
		11303-B16	
		27-65-B1	
		25065-B2	
	C204	15669-B1	
	E1	15597-B1	
	f1**	21816-B1	
	f2**	23724-B9	
	FCZ	15593-B1	
	fd**	25404-B1	
		29746-B1	
		23631-B1	
		25868-B1	
		25298-B1	
		25298-B2	
		11303-B37	
		11303-B24	
		11303-B26	
	If1**	11303-B27	
		11303-B28	
		11303-B29	
		11303-B30	
		11303-B33	
		11303-B31	
		11303-B25	
		11303-B35	
		11303-B34	
	MS2**	11303-B36	
	MU9	11303-B32	
	Mu-1	13706-B5	
	Ox6	11303-B1	
	P1**	11303-B2	
	P4 sid, **	11303-B3	
	Q-β**	11303-B4	
	R17**	35060-B1	
	Z1K/1	35060-B2	
	ZJ/2	35060-B3	
		11303-B5	
		11303-B6	
		11303-B7	
		11303-B38	
		12141-B1	

<i>Escherichia coli</i> (Cont'd)		11303-B20	The American Type Culture Collection
		11303-B17	
		11303-B15	
		11303-B11	
	547	11303-B18	
	UV1	13706-B2	
	UV47	23724-B2	
	UV375	23724-B1	
	$\alpha 3^{**}$	23724-B3	
	λ^{**}	23724-B4	
	λ C-17	23724-B5	
	λ sus P-3	23724-B6	
	λ sus R-5	23724-B7	
	λ sus J-6	23724-B8	
	λ sus O-8	35860-B1	
	λ sus A-11	13706-B3	
	λ ind'	15597-B2	
	$\phi 92$	13706-B1	
	ϕR	49696-B1	
	$\phi V-1$		
	$\phi X174^{**}$		
	$\phi Xcs70am-3$		
	G4** & ϕK^{**}		Biochim.Biophysica Acta.1992.1130:277-288
	BF23**		J.Bacteriol.1977.129:265-275
	Mu1		J.Ultrastruct.Res.1966.14:441-448
	Hp17		J.Mol.Biol.1991.218:705-721
	K3** & Ox2**		FEBS Lett.1987.215:145-150
	Rb18**, Rb51 & Rb69**		J.Bacteriol.1990.172:180-186
	H1**, H3, H8, K9, K18 & Ox1		Mol.Gen.Genet.1990.221:491-494
	M1**, Tula** & Tulb**		J.Mol.Biol.1987.196:165-174
	K10		J.Bacteriol.1979.140:680-686
	Qsr'		J.Bacteriol.1985.162:256-262
	B278		J.Gen.Microbiol.1988.134:1333-1338
	ϕ i 80**		FEMS Microbiol.Lett.1994.119:71-76
	ϕ i m173		Genetika 1985.21:673-675
	tf-1		J.Gen.Microbiol.1987.133:953-960
	P4 & ϕ iR73		Mol.Microbiol.1995.18:201-208
	I ₂ -2		J.Gen.Microbiol.1982.128:2797-2804
	PRD1		Virology 1990.177:445-451
	K3hx		Mol.Gen.Genet.1987.206:110-115
	933J** & 933W**		Infect.Immunity.1986.53:135-140
	H19-B**		J.Bacteriol.1987.169:4308-4312
	Tcp-111		Zentralbnl.Bakteriol.Mikrobiol.Hyg.1988.270:41-51

<i>Escherichia coli</i> (Cont'd)	N4**	Vet.Microbiol.1992.30:203-212
	Phi 80 trp	Ann.Inst.Pasteur.1971.120:121-125
	Obeta 1	J.Bacteriol.1978.133:172-177
	P1CM	J.Gen.Microbiol.1978.107:73-83
	PA-2**	J.Bacteriol.1990.172:1660-1662
	186**	Mol.Gen.Genet.1982.187:87-95
	186.IX.B	Mol.Microbiol.1992.6:2629-2642
	21**	Virology 1983.129:484-489
	P4**	MicrobiolRev.1993.57:683-702
	82**	J.Biol.Chem.1987.262:11721-11725
	PSP3	J.Bacteriol.1996.178:5668-5675
	HK022**	Nucleic Acids Res.1994.22:354-356
	D108**	Nucleic Acids Res.1986.14:3813-3825
	Rb49	J.Mol.Biol.1997.267:237-249
	Ike**	J.Mol.Biol.1985.181:27-39
	P22dis	Mol.Gen.Genet.1978.166:233-243
	N15**	J.Bacteriol.1996.178:1484-1486
	If1**	Proc.R.Soc.Lond.B.Biol.Sci.1991.245:23-30
	Stx2Phi-I & Stx2Phi-II	Infect.Immun.1998.66:4100-4107
	18	Virology 1987.156:122-126
	X	J.Gen.Microbiol.1981.126:389-396
	AC3	Mol.Microbiol.1991.5:715-725

	BW-1 C-1 E920g Esc-7-11 H19J Haiti HK243 Ia K20 K30 KL ₃ M Mu** O103 O157:H7 PID pt1 PilH α PR64FS PR772 SS4 β 4Q λ vir** Ω 8 09-1 92		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Haemophilus influenzae</i>	HP1**		Nucleic Acids Res. 1996.24:2360-2368
	S2**		Gene 1997. 196: 139-144
<i>Halobacterium cutirubrum</i>	S45		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Halobacterium halobium</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
			Can.J.Microbiol.1982.28:916-921
<i>Halobacterium salinarium</i>			Biol.Chem.Hoppe Seyler 1994.375:747-757

<i>Klebsiella oxytoca</i>	tf-1		J.Gen.Microbiol.1987.133:953-960
<i>Klebsiella pneumoniae</i>	60	23356-B1	The American Type Culture Collection
	92	23357-B1	
	K19Q		
	FC3-1 & FC3-9		
	FC3-10		FEMS Microbiol.Lett.1991.67:291-297
<i>Klebsiella sp.</i>	K11**		Mol.Gen.Genet. 1990.221:283-286
<i>Leptospira sp.</i>	LE1, LE3 & LE4		Res.Microbiol.1990.141:1131-1138
<i>Listeria monocytogenes</i>	243	23074-B1	The American Type Culture Collection
	197,1313 & 9425		Appl.Environ.Microbiol.1997.63:3374-3377
	H387 & H387-A		Appl.Environ.Microbiol.1993.59:2914-2917
	5775,6223 & 12682		APMIS.1993.101:160-167
	2389, 2671, 4211 & 2685		Intervirology 1994.37:31-35 & Zentralbl.Bakteriol.Mikrobiol.Hyg.1986.261:12-28
	4b, 4ab, 4g & 3c		Ann.Microbiol (Paris) 1977.128:185-198
	A118, A500 & A511**		Mol.Microbiol. 1995.16:1231-1241-992
	1, 3, 4, 5, 6, 7, 8, 9, 10, 11, 14, 15, 16, 17, 19 & 20		Ann.Microbiol. (Paris) 1979.130B:179-189
	1/2a, 1/2b, 3c, 4ab, 6a & 6b		Clin.Invest.Med.1984.7:229-232
	φLMUP35 2685		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Listeria innocua</i>	4211		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Micrococcus luteus</i>		4698-B1	The American Type Culture Collection
		4698-B4	
	N3	4698-2	
	N4	4698-B3	
	N8		
<i>Micrococcus luteus</i>	N17		Can.J.Microbiol. 1979.25:1027-1035
<i>Mycobacterium smegmatis</i>	BK-3	27203-B1	The American Type Culture Collection
	Bo1**	27204-B1	
	Bo 6	27205-B1	
	Bo 6II	27205-B2	
	Bo 6III	27205-B3	
	Mc-2	607-B6	
	Mc-4	607-B7	
	NN	11727-B1	
	Phagus lacticola	11759-B1	
	R1	607-B1	

	HER 317 HER 330 HER 333 HER 335 HER 334 HER 331 HER 316	Felix d'Herelle Refrence Centre, Quebec, Quebec
Legendre Leo Roy Sedge		
		Mol.Microbiol.1993.7:395-405
		J.Mol.Biol.1998.279:143-164
		Proc.Natl.Acad.Sci USA.1988.84:2833-2837
		Mol.Biol.Rep. 1981.30:11-15
		Proc.Natl.Acad.Sci.USA 1997.94:10961-10966
	29M, 31M, 122, 154, 37, 29D, 46, 139, 110, 141, 74D, AG1 & DS6A	Arch.Virol.1993.133:39-49 & Am.Rev.Respir.Dis.1975.112:17-22
<i>Mycobacterium fortuitum</i>	Bo 4 Bo 7	23052-B1 27207-B1 27207-B2 The American Type Culture Collection

<i>Mycobacterium leprae</i>			Ann.Microbiol. (Paris) 1982.133:93-97
<i>Mycobacterium tuberculosis</i>	DS6A	25618-B1 25618-B2 4243-B1	The American Type Culture Collection
	110, 139 & 33D		Arch.Virol.1993.133:39-49
	AG1,GS4E, BG1, PH & BK1		The Biology of Mycobacteria.Academic Press,Toronto 1982 (Ratledge & Stanford) 1982.309-351
<i>Mycobacterium sp</i>	Phagus pellegrini NN B1	11760-B1 11761-B1 23239-B1	The American Type Collection Culture

	TM4, ph60, ph72, PhAE39, phAE40 & Bxb1		Microbiology 1995.141:1173-1181
	C2		Experientia 1969.25:1112-1113
	18 & I15		J.Gen.Virol.1987.68:949-956
	63		Gruzlica 1968.36:617-622
	phlei & butyricum		J.Gen.Virol.1975.29:235-238
	MyF3P-59a		Z.Allg.Mikrobiol.1968.8:29-37
	Bo2a		J.Gen.Virol.1973.20:75-87
	D4,D28 & D32		J.Exptl.Med.1966.123:327-340
	HC		J.Bacteriol.1963.86:608-609
<i>Mycobacterium vaccae</i>	B5	15483-B1	The American Type Culture Collection
<i>Mycobacterium phlei</i>	NN Bo 2 Bo 2h Bo 3	11728-B1 11758-B1 27086-B2 27086-B1	The American Type Culture Collection
<i>Mycoplasma arthritidis</i>	MAV1**		Infect.Immunity.1995.63:4016-4023
<i>Mycoplasma hyorhinis</i>	Hr-1		Arch.Virol.1983.77:81-85
<i>Mycoplasma pneumoniae</i>	Br-1		Arch.Virol.1983.75:1-15
<i>Mycoplasma pulmonis</i>			Plasmid 1995. 33: 41-49
<i>Mycoplasma sp.</i>			J.Gen.Microbiol.1985:131:3117-3126
			J. Virol.1986.59:584-590
			Gene 1994. 141: 1-8

		Microbios 1990. 64: 111-125
		Infection & Immunity 1995. 63: 4016-4023
		Med.Biol.1982.60:116-120
MV-L2 &		Arch.Virol.1979.61:289-296
		Acta.Virol.1978.22:443-450
		J.Gen.Virol.1979.42:315-322
		Virology 1973.55:118-126

			Science 1971.173:725-727
<i>Neisseria perflava</i>			J.Clin.Microbiol.1976. 4:87-91
<i>Nocardia erythropolis</i>	φC		J.Gen.Virol.1974.23:247-254
	φEC		J.Bacteriol.1976.126:1104-1107
<i>Pasteurella multocida</i>	B225		Arch.Exp.Veterinarmed.1981.35:433-436
	B939a		Am.J.Vet.Res.1978.39:1565-1566
	Nos.115, 32, 967 & 1075		Vet.Med.Nauki. 1977.14:33-36
<i>Propionibacterium acnes</i>	NN	29399-B1	The American Type Collection Culture

<i>Pseudomonas aeruginosa</i>	2	12175-B1	The American Type Culture Collection
	2A	12175-B2	
	2B	12175-B3	
	11	12175-B4	
	16	14205-B1	
	24	14206-B1	
	27	14207-B1	
	44	14208-B1	
	73	14209-B1	
	95	14210-B1	
	109	14211-B1	
	113	14212-B1	
	249	14213-B1	
	B3	14214-B1	
	Hoff 2	15692-B1	
	Hoff 3	14203-B1	
	Pa	14204-B1	
	Pb	12055-B1	
	PB-1	12055-B2	
	Pc	15692-B3	
	Pf	12055-B3	
	PP7**	25102-B1	
		15692-B2	
	7 & 31		Felix d'Herelle Reference Centre, Quebec, Quebec
	PF3**		J.Virol.1983.47:221-223
	φ-MC		Can.J.Microbiol.1969.15:1179-1186
	PF1**		J.Mol.Biol.1991.218:349-364
	PR4**		J.Gen.Virol.1979.43:583-592
	A7		J.Bacteriol.1992.174:2407-2411
	KF1		J.Biochem.1983.93:61-71
	φCTX**		Mol.Microbiol.1993.4:1703-1709
	φ2**		J.Virol.1977.24:135-141

	<p>φKZ, 21, φNZ, PMN17, PTB80, 68, PB-1, E79, 16, 109, 352, 1214, F8, 71, 337, M4, φC17, SL2, B17, Li-24, φmnP78, PS17**, φ1, 73, M6, Li-2, 7, φmnF82, PTB2, PTB20, PTB42, φKF77, 31, PTB21, 119x, φPLS27, B3, 258, Hw12, PM57, PM62, PM105, 148, PM681, 198, 218, 222, 242, 246, PC131, φC11, SL5, D3112**, Jb19, F7, PM69, PM13, PM61, PM113, φ240, 249 & 269</p>		ddd
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<i>Pseudomonas aeruginosa</i> (Cont'd)	297, 309, 318, 11,		Arch. Virol. 1993. 131:141-151
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<i>Pseudomonas cepacia</i>			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Pseudomonas fragi</i>		27362-B1 27363 B1	The American Type Culture Collection
	wy		
<i>Pseudomonas phaseolicola</i>	φ6		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Pseudomonas putida</i>	gh-1	12633-B1	The American Type Culture Collection
<i>Pseudomonas syringae</i>		40492-B1 21781-B1	The American Type Culture Collection
	φ6		
<i>Pseudomonas sp.</i>	PPs-G3	49780-B1	The American Type Culture Collection
<i>Salmonella bareilly</i>	Sab 2		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Salmonella enteritidis</i>	1, 2, 3 & 6		Epidemiol. Infect. 1995.114:227-236
	2a, 3a, 4a, 5a, 6a, 7a, 8a, 9a, 15, 19, 20 & 21**		Vet. Med. Nauki. 1975.12:55-60
<i>Salmonella newington</i>	Epsilon 34		J. Struct. Biol. 1995.115:283-289
<i>Salmonella newport</i>		27869-B1 27869-B2	The American Type Culture Collection
	16-19		
			Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Salmonella paratyphi</i>		19940-B1 12176-B1	The American Type Culture Collection
	Paratyphoid A		
	Jersey		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Salmonella senftenberg</i>	SasL1, SaL2, Sal 3, SaL4, SaL5 & SasL6		Indian J. Med. Res. 1997.105:47-52
<i>Salmonella typhimurium</i>	P22**	19585-B1	The American Type Culture Collection
	SL-1	40282	
	MB78**		J. Virol. 1982.41: 1038-1043
	SE1		J. Gen. Microbiol. 1986.132:1035-1041
	LT2		Virology 1971.45:835-636
	ES18**		Virology 1970.42:621-632
	L**		J. Virol. 1985.56:1034-1036

	P1CM clr-100		Mol.Gen.Genet.1975.138:113-126
	F22		Genet.Res.1986.48:139-143
	Fels 1		J.Gen.Virol.1978.38:263-272
	Fels 2		Genet.Res.1986.48:139-143
	Px		Mol.Gen.Genet.1970.108:184-202
	P1kc		Virology 1974.60:503-514
	A3 & A4		J.Bacteriol. 1987.169:1003-1009
	HT		Genet.Res.1976.27:315-322
<i>Salmonella typhimurium</i> (Cont'd)	IRA		J.Basic Microbiol. 1990.30:707-716
	Mud1		Mol.Gen.Genet. 1986.202:327-330
	P22 (cir4-1, cir5-1 & cir6-1)		Mol.Gen.Genet.1984.198:105-109
	BF23**		Mol.Gen.Genet.1976.147:195-202
	Kb1		J.Bacteriol.1974.117:907-908
	P221dis		J.Gen.Virol.1978.41:367-376
	PRD1**		Virology 1990.177:445-451
	I ₂ -2**		J.Gen.Microbiol.1982.128:2797-2804
	tf-1		J.Gen.Microbiol.1987.133:953-960
	X**		J.Gen.Microbiol.1981.126:389-396
<i>Salmonella typhosa/typhi</i>	8	19937-B1	The American Type Culture Collection
	23	19938-B1	
	25	19939-B1	
	46	19942-B1	
	53	19943-B1	
	163	19946-B1	
	175	19947-B1	
	Vil	27870-B1	
	ViVI	27870-B2	Felix d'Herelle Refrence Centre,Quebec,Quebec
	O1		
	VIII		
	j2		
<i>Salmonella sp.</i>	P3	25957-B1	The American Type Culture Collection
	P4**	25957-B2	
	P9a	25957-B3	
	P9c	25957-B4	
	P10	25957-B5	
	102	19945-B1	
	Chi (χ)	9842-B1	
	R34	97541	
	MG40		Virology 1968.34:521-530
	P14		Microb.Pathog.1990.8:393-402
	PSP3		Virology 1992.188:414
	Ike**		Zentralbl.Bakteriol.1976.234:294-304
<i>Sphaerotilus natans</i>	P27 & 9NA		J.Virol.1986.12:921-931
	SN1		Appl.Environ.Microbiol.1979.37:1025-1030

<i>Shigella dysenteriae</i>	P2 φ80	23351-B1 11456b 11456a-B1	The American Type Culture Collection
<i>Shigella flexeneri</i>	D20	12661-B1	The American Type Culture Collection
	SfII**		Mol.Microbiol.1997.26:939-950
	SfV**		Gene 1997.22:217-227
	Sf6**		Mol.Microbiol.1995.18:201-208
	SfX		Gene 1993.129:99-101
<i>Shigella sonnei</i>	C16**		
	Ufa		Mol..Biol (Mosk) 1977.11:323-331
<i>Shigella sp</i>	37	23354-B1	The American Type Culture Collection
<i>Spiroplasma citri</i>	SpV1		Plasmid 1993.29:193-205
<i>Spiroplasma sp.</i>	SpV1-R8A2B		Nucleic Acids Res. 1990.18:1293
	SpV3		Isr.J.Med.Sci.1987.23:429-433
	Sp V4		J.Bacteriol.1987.169:4950-4961
<i>Staphylococcus albus</i>			Staphylococci & Staphylococcal Infections.1997. Voll:503-508 (Karger,Basel)

<i>Staphylococcus aureus</i>			The American Type Culture Collection
		27702-B1	
		27703-B1	
		27704-B1	
		23360-B1	
		23361-B1	
	15	27705-B1	
	17	27712-B1	
	29	27690-B1	
	42D**	27691-B1	
	42E	27692-B1	
	47	27693-B1	
	52	27694-B1	
	52A	27695-B1	
	53	27696-B1	
	54	27697-B1	
	55	27698-B1	
	71	27699-B1	
	75	27693-B2	
	77	27700-B1	
	79	27701-B1	
	80	27706-B1	
	81	27707-B1	
	83A	27708-B1	
	84	33742	
	85**	33741-B1	
	88	15565	
	92	19685-B1	
	5504'	11987-B1	
	K	11988-B1	
	P1	15752-B1	
	P14		
	UC18		

	HER 101 HER 239 HER 283 HER 49	Felix d'Herelle Reference Centre, Quebec, Quebec
Twort**		
$\phi 11^{**}$		J.Bacteriol.1988.170:2409-2411
$\phi 13^{**}$ & $\phi 42^{**}$		J.Gen..Microbiol.1989.135:1679-1697
L54a**		J.Bacteriol.1986.166:385-391
80 α^{**}		Can.J.Microbiol.1996.43:612-616
94,95 & 96		J.Clin.Microbiol.1988.26:2395-2401
$\phi 131, A_3$ & A_5		Staphylococci & Staphylococcal Infections.1997. Vol1:503-508 (Karger,Basel)
Phi PVL**		Gene 1998.215:57-67
<i>Staphylococcus carnosus</i>	BaSTC2	Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Staphylococcus epidermidis</i>	1a, 2b, 3a, 4b, 5a, 6b, 7b, 8c, 9a, 10a, 11b, 12a & 13b	Can.J.Microbiol.1988.34:1358-1361
	41, 63, 118II, 138, 245, 336, 392 & 550	Res.Virol.1994.145:111-121
<i>Staphylococcus saprophyticus</i>	1154A, 1405, 1314, 1139 & 1259	Res.Virol.1990.141: 625-635 & Res.Virol.1994.145:111-121
<i>Staphylococcus sp.</i>	Phi 812, Phi 131, SK311 & U16	Virology 1998.246:241-252
<i>Streptococcus faecalis</i>	VD13	HER44 Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Streptococcus faecium</i>	PE1	Zentralbl.Bakteriol.1975.231:421-425
<i>Streptococcus oralis</i>	Cp-1** & Cp- 7**	FEMS Microbiol.Lett.1989.65:187-192

<i>Streptococcus pneumoniae</i>	Cp-1**	HER223	Felix d'Herelle Reference Centre, Quebec, Quebec
	Cp-1**, Cp-5**, Cp-7**, Cp-9**, ω-1 & ω-2		J.Virol.1981.40:551-559 & Eur.J.Biochem.1979.101:59-64 & Microbial Drug Resistance 1997.3:165-176
	HB-623 & HB-746		J.Virol.1990.64:5149-5155
	EJ-1**		J.Bacteriol.1992.174:5516-5525
	Dp-2 & Dp-4		J.Virol.1978.26:221-225
	Dp-1		Virology 1975.63:577-582
	ω-3 & ω-8		J.Virol.1976.19:659-667
	304		J.Bacteriol.1980.141:1298-1304
	HB-1, HB-2, HB-3**, HB-4, HB-5 & HB-6		J.Bacteriol.1979.138:618-624
<i>Streptococcus pyogenes</i>	T12**		Mol. Microbiology. 1997#23:719-728
	A-1 A-6 A-25 Kjem	12202-B1 12203-B1 12204-B1 14918	The American Type Culture Collection
	1 182 VD1884	HER 339 HER 80 HER 323	Felix d'Herelle Reference Centre, Quebec, Quebec
	1A 1B NN 42 118 120	12169-B1 12170-B1 21597-B1 19948-B1 19951-B2 19952-B1	The American Type Culture Collection
<i>Veillonella rodentium</i>	N2		Antonie Van Leeuwenhoek 1989.56:263-271
<i>Vibrio cholerae</i>	Psi 92		Intervirology 1993.36:237-244
	VCB-1,2,3 & 4		J.Infection 1998.36:131
	CP-T1**		J.Virol.1984.51:163-169
	VSK		FEMS Microbiol.Lett.1996.145:17-22
	Phil38		J.Virol.1986.57:960-967
	Phil49		J.Virol.1985.140:217-223
	Fs-2**		Microbiology 1998.144:1901-1906

	e4 e5 X29 β κ 13 14 16 24 32 57		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio cholerae</i> (Cont'd)	138 145 149 163 N-4 S-5 S-20 M-4 D-10 I II III IV V	14100-B1 14100-B2 14100-B30 14100-B4 51352-B1 51352-B2 51352-B3 51352-B4 51352-B5 51352-b6 51352-B7 51352-B8 51352-B9 51352-B10	The American Type Culture Collection
<i>Vibrio costicola</i>	UTAK		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio eltor</i>	e ₄		J.Gen.Virol.1987.68:1411-1416
<i>Vibrio natrigens</i>	nt1, nt6		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio</i> <i>parahaemolyticus</i>	KVP40** VF33 VP1 ϕ 60 ϕ HAWI-5 ϕ PEL8C-1		Felix d'Herelle Reference Centre, Quebec, Quebec
<i>Vibrio</i> sp.	α 3a		Felix d'Herelle Reference Centre, Quebec, Quebec
	NN ph1	11985-B1 51582-B1	The American Type Culture Collection
	Phi149		J.Virol.1987.61:3999-4006
<i>Veillonella rodentium</i>	N2		Antonie V.Leeuwenhoek.1989.56:263-271

<i>Yersinia enterocolitica</i>	1 2 3 4 5 6 7 8 9 ϕ YeO3-12		Felix d'Herelle Reference Centre, Quebec, Quebec
	I, IV & VIII		Zentralbl.Bakteriol.Mikrobiol.Hyg.1982.253:1 02
<i>Yersinia pestis</i>	R S Y	23208-B1 11593-B1 23053-B1	The American Type Culture Collection
	II		Zh.Mikrobiol.Epidemiol.Immunobiol.1990.11 :9
<i>Yersinia pseudotuberculosis</i>	PST**	23207-B1	The American Type Culture Collection
<i>Yersinia sp.</i>	RD2		Mol.Gen.Mikrobiol.Virusol.1990.8:18-21

xxxx)

Table 2

>Bacteriophage 77, complete genome sequence, 41708 nucleotides

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1   gatcaaaata cttggggaac ggtagggag taaacttcgc gataatttta aaaattcatg
61  tataaccccc ctcttataac cattttaagg caggatgatg aatggagatt atagtcgatg
121 aaaatttagt gcttaaagaa aaagaaaggc tacaagtatt atataaagac atacctagca
181 ataaattaaa agtagttgat gggttaatta ttcaagcagc aaggctacgt gtaatgcctg
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39181	atggatgggt	tcgacatctt	tattgttggg	atactgtcat	tattcggtat	attcgcatgt
39241	ctacttgTTa	tcacattgcc	tatctataca	gtggctagtt	accaacacaa	agaattacat
39301	caaggaaacta	ttacagataa	atataacaag	agacaagata	aagaagacaa	gttctatatt
39361	gtattagaca	acaaacaagt	cattgaaaat	tccgacttat	tattcaaaaa	gaaatttgat
39421	agcgcagata	tacaagctag	gttaaaaagta	ggcgataaagg	tagaagttaa	aacaatcggg
39481	tatagaatac	acttttttaa	tttatatccg	gtcttatacg	aagtaaagaa	ggtagataaa
39541	caatgattaa	acaaatacta	agactattat	tcttactagc	aatgtatgag	ttaggttaagt
39601	atgtaactga	gcaagtgtat	attatgatga	cggctaataga	tgatgtagag	gcgccgagtg
39661	attactgctt	tcgagcggag	gtgagtgaat	aatgagaata	tttatttatg	atttgatcgt
39721	tttgctgTTt	gtcttcttaa	tatccatata	tattattgat	gatggagtga	taataaatgc
39781	attaggaatt	tttggtatgt	ataaaaattat	agattccttt	tcagaaaata	ttataaagag
39841	gtagataaaa	atgaacgagc	aaataatagg	aagcatatat	acttttagcag	gaggtgttgt
39901	gctttattca	gttaaagaga	tttttaggta	ttttacagat	tctaacttac	aacgtaaaaa
39961	aatcaattta	gaacaaatat	atccgatata	tttagattgt	tttaaaaagg	ctaaaagat
40021	gattggagct	tatatatttc	caacagaaca	gcatgaattt	ttagattttt	ttgatattga
40081	agtctttaat	aatttagata	agcaaagtaa	aaaagcgtat	gaaaatgtta	ttggatttag
40141	acaaatgatt	aatttatcaa	atagagttaa	ggcaatggaa	gattttaaga	tgagtttcaa
40201	caatgaattt	agtacaaatc	agattttttt	taatccttct	tttgttatgg	aaacaattgc
40261	tattataaat	gaatatcaaa	aagatatatc	ttatttaaaa	aatataatta	ataaaatgaa

40321 tgaaaataga gcttataatc atattgatag ttttatcact tcagagtacc gacgaaaaat
40381 aaacgattat aatctttatc ttgataaatt tgaagaacag tttagtcaaa agtttaaaat
40441 aaacagaact tcgataaaaag aaagaattat tattaattta aacaagagga gatttaaatg
40501 atgtggatta ctatgactat tgtatttgct atattgctat tagtttgat cagtattaat
40561 agtgcctg caagagagat acaagcactt agatatatga atgattatct acctgatgaa
40621 gtagttaaaa ctaaagggtta caacgggtta gaagaatata ggattgaatt gaagcgaatg
40681 aataacgata ttaaaaagta atttatatta tcggagggtat tgcattgaat gataaagatt
40741 gagaaacacg atatcaaaaa gcttgaagaa tacattcagc acatcgataa ctatcgaaga
40801 gagttgaaga tgcgagaata tgaattactt gaaagtcag aaccagataa tgcgggagct
40861 ggcaaaagta atttgccggg taacccgatt gaacgatgtg caataaagaa gtttagtgat
40921 aacaggtaaca atacattaag aaatatagtt aacgggtgtag atagattgat aggtgaaagt
40981 gatgaggata cgcttgagtt attaaggttt agatattggg attgtcctat tggttggtat
41041 gaatgggaag atatagcaca ttactttggt acaagtaaga caagtatat acgtagaagg
41101 aatgcactga tcgataagtt agcaaagtat attggttatg tgtagcggac ttttacccta
41161 tgtaagtccg cattaaaaca gtttattatg ttagtatcag attaatattt aaagttatta
41221 aatgctaata cgacgcatga acaagaggcg catcactatg tgatgtgctt ttttatttat
41281 gaggtatgaa catgttcaaa ctaattgtaa atacattact acacatcaag tatagatgag
41341 tcttgatact acttaagtta tataagggtga aacattatga tgactaaaga cgaacgtata
41401 cgattctata agtctaaaga atggcaaata acaagaaaaa gagtgctaga aagagataat
41461 tatgaatgtc aacaatgtaa gagagacggc aagttaacga catatgacaa aagcaagcgt
41521 aagtcggttg atgtagatca tatattatcg ctagaacatc atccggagtt tgctcatgac
41581 ttaacaatt tagaaacact gtgtattaaa tgtcacaaca aaaaagaaaa gagatttata
41641 aaaaagaaa ataaatggaa agacgaaaaa tggtaaatat ccccggttca aaaaaatcaa
41701 aagcgatc

Table 3

	Name	Position		Name	Position
1	77ORF005	19572..21026	48	77ORF052	1762..2013
2	77ORF006	3976..5196	49	77ORF053	37521..37757
3	77ORF007	21871..23076	50	77ORF054	22818..23060
4	77ORF008	2120..3307	51	77ORF055	17546..17788
5	77ORF009	31946..32803	52	77ORF058	18892..19122
6	77ORF010	26092..26889	53	77ORF059	34564..34785
7	77ORF011	24441..25208	54	77ORF064	29574..29795
8	77ORF012	29788..30576	55	77ORF065	28528..28746
9	77ORF013	33620..34399	56	77ORF066	27494..27703
10	77ORF014	27760..28512	57	77ORF069	38341..38547
11	77ORF015	3291..4028	58	77ORF070	36269..36475
12	77ORF016	32867..33610	59	77ORF071	40498..40701
13	77ORF017	23269..23982	60	77ORF072	38735..38938
14	77ORF018	31169..31840	61	77ORF073	30945..31148
15	77ORF019	39851..40501	62	77ORF074	38544..38738
16	77ORF020	6926..7570	63	77ORF075	13673..13870
17	77ORF021	37762..38304	64	77ORF077	25357..25605
18	77ORF022	30605..31156	65	77ORF079	29089..29280
19	77ORF023	26903..27346	66	77ORF080	35204..35389
20	77ORF024	10700..11140	67	77ORF085	24060..24242
21	77ORF025	9707..10147	68	77ORF092	39706..39876
22	77ORF026	40729..41145	69	77ORF094	32226..32393
23	77ORF027	6518..6925	70	77ORF096	13606..13773
24	77ORF028	34795..35199	71	77ORF098	7092..7256
25	77ORF029	6117..6521	72	77ORF102	29051..29212
26	77ORF030	36478..36879	73	77ORF104	34393..34551
27	77ORF031	39151..39546	74	77ORF109	18282..18434
28	77ORF032	33892..34266	75	77ORF112	39543..39692
29	77ORF033	5758..6120	76	77ORF117	27361..27501
30	77ORF034	7886..8236	77	77ORF118	38390..38530
31	77ORF035	19258..19560	78	77ORF120	36059..36199
32	77ORF036	36876..37223	79	77ORF124	33699..33833
33	77ORF037	102..446	80	77ORF128	14221..14355
34	77ORF038	34908..35219	81	77ORF130	15675..15806
35	77ORF039	37220..37528	82	77ORF133	8414..8542
36	77ORF040	41377..41676	83	77ORF140	13113..13235
37	77ORF041	35454..35753	84	77ORF147	7029..7148
38	77ORF042	5490..5774	85	77ORF149	30668..30787
39	77ORF043	29304..29564	86	77ORF151	31837..31953
40	77ORF044	18481..18768	87	77ORF155	30278..30391
41	77ORF045	5216..5500	88	77ORF157	4044..4157
42	77ORF046	25663..25935	89	77ORF167	20692..20799
43	77ORF047	11159..11425	90	77ORF175	35717..35821
44	77ORF048	28776..29039	91	77ORF176	6836..6940
45	77ORF049	36013..36255	92	77ORF178	35390..35491
46	77ORF050	35753..36007	93	77ORF179	8318..8419
47	77ORF051	38931..39167	94	77ORF182	29268..29564

Table 4

77ORF017 sequence

```

23982      atgacgcataatatagaaaaacgcattaataaattaaaaacttct
1      M  T  H  N  I  E  K  R  I  N  K  L  K  T  S
23937      ggaaatccaaaattttaaaaagtttagattcagatattcactattta
16     G  N  P  K  F  K  K  L  D  S  D  I  H  Y  L
23892      ctcaagagattttgaaggtgaaaaaaaccataaagggtttttatcca
31     L  K  R  F  E  G  E  K  N  H  K  G  F  Y  P
23847      aagtttaacaaggagaaatagttttttagatttcggtataaac
46     K  F  K  Q  G  E  I  V  F  V  D  F  G  I  N
23802      gttaataaagaatttttctaattcacactttgcaatagtgatgaat
61     V  N  K  E  F  S  N  S  H  F  A  I  V  M  N
23757      aaaaatgattctaataacggaggatatagtaaatgttattccctta
76     K  N  D  S  N  T  E  D  I  V  N  V  I  P  L
23712      tcctctaagaaaaacaaaaagtatttaagatgaattttgatttg
91     S  S  K  E  N  K  K  Y  L  K  M  N  F  D  L
23667      aaatgggagtattatttaagattgtttttaaatttaattagcgcg
106    K  W  E  Y  Y  L  R  L  F  L  N  L  I  S  A
23622      caaaataattcagctatatataaaagaagttttcgataaaaaaatac
121    Q  N  N  S  A  I  L  K  E  V  F  D  K  K  Y
23577      caaaaaaacaacacagaattcatcactaaagattattttattgaa
136    Q  K  N  N  T  E  F  I  T  K  D  Y  F  I  E
23532      tttatatctgatagtttagaaattgaaaataaattaaataaaatt
151    F  I  S  D  S  L  E  I  E  N  K  L  N  K  I
23487      gacagaaacattaataacatagtatcagcaattgataaggtaaaa
166    D  R  N  I  N  N  I  V  S  A  I  D  K  V  K
23442      aaattaaaaggtaatatgttacgcttgcataaattctttccagccg
181    K  L  K  G  N  S  Y  A  C  I  N  S  F  Q  P
23397      attagtaagtttcgcataagaaaagttttacccccaaaaaattaaa
196    I  S  K  F  R  I  R  K  V  L  P  Q  K  I  K
23352      aatccagtaatatagattcttcggatattatgttactgataaataga
211    N  P  V  I  D  S  S  D  I  M  L  L  I  N  R
23307      attaataataatatattgcagatccctgatataagatga 23269
226    I  N  N  N  I  L  Q  I  P  D  I  R  *

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Physico-chemical parameters of ORF 77ORF017

1 MTHNIEKRIN KLKTSGNPKF KKLDSDIHYL LKRFEGEKNH KGFYPKFKQG EIVFVDFGIN
 61 VNKEFSNSHF AIVMNKNSDN TEDIVNVIPL SSKENKKYLK MNFDLKWEYY LRLFLNLISA
 121 QNNSAILKEV FDKKYQKNNT EFITKDYFIE FISDSLEIEN KLNKIDRNIN NIVSAIDKVK
 181 KLKGSYACI NSFQISKFR IRKVLQKIK NPVIDSSDIM LLINRINNNI LQIPDIR

Number of amino acids: 237
 Average molecular weight (Daltons): 27887.38
 Mean amino acid weight (Daltons): 117.67
 Monoisotopic molecular weight (Daltons): 27869.83
 Mean amino acid monoisotopic weight (Daltons): 117.59

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	5	2.11%	7.58%	Cys	C	1	0.42%	1.66%
Asp	D	14	5.91%	5.28%	Glu	E	13	5.49%	6.37%
Phe	F	16	6.75%	4.09%	Gly	G	6	2.53%	6.84%
His	H	4	1.69%	2.24%	Ile	I	29	12.24 %	5.81%
Lys	K	33	13.92 %	5.95%	Leu	L	19	8.02%	9.42%
Met	M	4	1.69%	2.37%	Asn	N	30	12.66 %	4.45%
Pro	P	7	2.95%	4.9%	Gln	Q	6	2.53%	3.97%
Arg	R	8	3.38%	5.16%	Ser	S	17	7.17%	7.12%
Thr	T	5	2.11%	5.67%	Val	V	11	4.64%	6.58%
Trp	W	1	0.42%	1.23%	Tyr	Y	8	3.38%	3.18%

Number of acidic (negative) amino acids (ED): 27
 11.39%
 Number of basic (positive) amino acids (KR): 41
 17.30%
 Total charge (KRED): 68
 28.69%
 Net charge (KR - ED): 14
 5.91%
 Theoretical pI: 10.01
 Total linear charge density: 0.30
 Average hydrophobicity: -5.37
 Ratio of hydrophilicity to hydrophobicity: 1.41
 Percentage of hydrophilic amino acid: 57.81%
 Percentage of hydrophobic amino acid: 42.19%
 Ratio of %hydrophilic to %hydrophobic: 1.37

77ORF019 sequence

```
39851      atgaacgagcaaataaataggaagcatatatacttttagcaggaggt
1   M   N   E   Q   I   I   G   S   I   Y   T   L   A   G   G
39896      gttgtgctttatttcagttaaagagatttttaggtattttacagat
16  V   V   L   Y   S   V   K   E   I   F   R   Y   F   T   D
39941      tctaacttacaacgtaaaaaaatcaatttagaacaatatatccg
31   S   N   L   Q   R   K   K   I   N   L   E   Q   I   Y   P
39986      atatatttagattgttttaaaaaggctaaaaagatgattggagct
46   I   Y   L   D   C   F   K   K   A   K   K   M   I   G   A
40031      tatattattccaacagaacagcatgaatttttagatttttttgat
61   Y   I   I   P   T   E   Q   H   E   F   L   D   F   F   D
40076      attgaagtctttaataatttagataagcaaagtaaaaaagcgtat
76   I   E   V   F   N   N   L   D   K   Q   S   K   K   A   Y
40121      gaaaatgttattggatttagacaaatgattaatttatcaaataga
91   E   N   V   I   G   F   R   Q   M   I   N   L   S   N   R
40166      gttaaggcaatggaagattttaagatgagtttcaacaatgaattt
106  V   K   A   M   E   D   F   K   M   S   F   N   N   E   F
40211      agtacaaatcagattttttttaatccttcttttgttatggaaaca
121  S   T   N   Q   I   F   F   N   P   S   F   V   M   E   T
40256      attgctattataaatgaatatcaaaaagatatatcttatttataaa
136  I   A   I   I   N   E   Y   Q   K   D   I   S   Y   L   K
40301      aatataattaataaaaatgaatgaaaatagagcttataatcatatt
151  N   I   I   N   K   M   N   E   N   R   A   Y   N   H   I
40346      gatagttttatcacttcagagtaccgacgaaaaataaacgattat
166  D   S   F   I   T   S   E   Y   R   R   K   I   N   D   Y
40391      aatctttatcttgataaatttgaagaacagtttagtcaaaagttt
181  N   L   Y   L   D   K   F   E   E   Q   F   S   Q   K   F
40436      aaaataaacagaacttcgataaaaagaagaattattattaattta
196  K   I   N   R   T   S   I   K   E   R   I   I   I   N   L
40481      aacaagaggagattttaaatga 40501
211  N   K   R   R   F   K   *
```

Physico-chemical parameters of ORF 77ORF019

1 MNEQIIGSIY TLAGGVVLYS VKEIFRYFTD SNLQRKKINL EQIYPIYLDL FKKAKKMIGA
 61 YIIPTEQHEF LDFFDIEVFN NLDKQSKKAY ENVIGFRQMI NLSNRVKAME DFKMSFNNEF
 121 STNQIFFNPS FVMETIAIIN EYQKDISYK NIINKMNENR AYNHIDSFIT SEYRRKINDY
 181 NLYLDKFEEQ FSQKFKNRT SIKERIIINL NKRRFK

Number of amino acids: 216
 Average molecular weight (Daltons): 26026.06
 Mean amino acid weight (Daltons): 120.49
 Monoisotopic molecular weight (Daltons): 26009.34
 Mean amino acid monoisotopic weight (Daltons): 120.41

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	7	3.24%	7.58%	Cys	C	1	0.46%	1.66%
Asp	D	10	4.63%	5.28%	Glu	E	16	7.41%	6.37%
Phe	F	19	8.80%	4.09%	Gly	G	5	2.31%	6.84%
His	H	2	0.93%	2.24%	Ile	I	28	12.96%	5.81%
Lys	K	22	10.19%	5.95%	Leu	L	12	5.56%	9.42%
Met	M	7	3.24%	2.37%	Asn	N	23	10.65%	4.45%
Pro	P	3	1.39%	4.9%	Gln	Q	10	4.63%	3.97%
Arg	R	11	5.09%	5.16%	Ser	S	13	6.02%	7.12%
Thr	T	7	3.24%	5.67%	Val	V	7	3.24%	6.58%
Trp	W	0	0.00%	1.23%	Tyr	Y	13	6.02%	3.18%

Number of acidic (negative) amino acids (ED): 26
 12.04%
 Number of basic (positive) amino acids (KR): 33
 15.28%
 Total charge (KRED): 59
 27.31%
 Net charge (KR - ED): 7
 3.24%
 Theoretical pI: 9.52
 Total linear charge density: 0.28
 Average hydrophobicity: -4.84
 Ratio of hydrophilicity to hydrophobicity: 1.37
 Percentage of hydrophilic amino acid: 54.17%
 Percentage of hydrophobic amino acid: 45.83%
 Ratio of %hydrophilic to %hydrophobic: 1.18

77ORF043 sequence

```
29304 .   atgtattacgaaataggcgaaatcatacgcaaaaatattcatgtt
1    M Y Y E I G E I I R K N I H V
29349     aacggattcgattttaagctattcatttttaaagggtcatatgggc
16   N G F D F K L F I L K G H M G
29394     atatcaatacaagttaaagatatgaacaacgtaccaattaaacat
31   I S I Q V K D M N N V P I K H
29439     gcttatgtcgtagatgagaatgacttagatatggcatcagactta
46   A Y V V D E N D L D M A S D L
29484     tttaaccaagcaatagatgaatggattgaagagaacacagacgaa
61   F N Q A I D E W I E E N T D E
29529     caggacagactaattaacttagtcatgaaatggtag 29564
76   Q D R L I N L V M K W *
```

Physico-chemical parameters of ORF 77ORF043

1 MYYEIGEIR KNIHVNGFDF KLFILKGHMG ISIQVKDMNN VPIKHAYVVD ENLDLMASDL
 61 FNQAIDEWIE ENTDEQDRLI NLVMKW

Number of amino acids: 86
 Average molecular weight (Daltons): 10186.68
 Mean amino acid weight (Daltons): 118.45
 Monoisotopic molecular weight (Daltons): 10180.02
 Mean amino acid monoisotopic weight (Daltons): 118.37

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.49%	7.58%	Cys	C	0	0.00%	1.66%
Asp	D	9	10.47%	5.28%	Glu	E	7	8.14%	6.37%
Phe	F	4	4.65%	4.09%	Gly	G	4	4.65%	6.84%
His	H	3	3.49%	2.24%	Ile	I	11	12.79%	5.81%
Lys	K	6	6.98%	5.95%	Leu	L	6	6.98%	9.42%
Met	M	5	5.81%	2.37%	Asn	N	8	9.30%	4.45%
Pro	P	1	1.16%	4.9%	Gln	Q	3	3.49%	3.97%
Arg	R	2	2.33%	5.16%	Ser	S	2	2.33%	7.12%
Thr	T	1	1.16%	5.67%	Val	V	6	6.98%	6.58%
Trp	W	2	2.33%	1.23%	Tyr	Y	3	3.49%	3.18%

Number of acidic (negative) amino acids (ED): 16
 18.60%
 Number of basic (positive) amino acids (KR): 8
 9.30%
 Total charge (KRED): 24
 27.91%
 Net charge (KR - ED): -8
 9.30%
 Theoretical pI: 4.38
 Total linear charge density: 0.30
 Average hydrophobicity: -2.80
 Ratio of hydrophilicity to hydrophobicity: 1.19
 Percentage of hydrophilic amino acid: 48.84%
 Percentage of hydrophobic amino acid: 51.16%
 Ratio of %hydrophilic to %hydrophobic: 0.95

77ORF102 sequence

```
29051      atgagcaacattttataaaagctacctagtagcagtattatgcttc
1      M  S  N  I  Y  K  S  Y  L  V  A  V  L  C  F
29096      acagtcttagcgattgtacttatgccgtttctatacttcactaca
16     T  V  L  A  I  V  L  M  P  F  L  Y  F  T  T
29141      gcatgggtcaattgcgggattcgcaagtatcgcaacattcatgtac
31     A  W  S  I  A  G  F  A  S  I  A  T  F  M  Y
29186      tacaaagaatgcttttttcaaagaataa 29212
46     Y  K  E  C  F  F  K  E  *
```

Physico-chemical parameters of ORF 77ORF102

1 MSNIYKSYLV AVLCFTVLAI VLMPLYFTT AWSIAGFASI ATFMYYKECF FKE

Number of amino acids: 53
Average molecular weight (Daltons): 6155.42
Mean amino acid weight (Daltons): 116.14
Monoisotopic molecular weight (Daltons): 6151.07
Mean amino acid monoisotopic weight (Daltons): 116.06

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	6	11.32 %	7.58%	Cys	C	2	3.77 %	1.66%
Asp	D	0	0.00%	5.28%	Glu	E	2	3.77 %	6.37%
Phe	F	7	13.21 %	4.09%	Gly	G	1	1.89 %	6.84%
His	H	0	0.00%	2.24%	Ile	I	4	7.55 %	5.81%
Lys	K	3	5.66%	5.95%	Leu	L	5	9.43 %	9.42%
Met	M	3	5.66%	2.37%	Asn	N	1	1.89 %	4.45%
Pro	P	1	1.89%	4.9%	Gln	Q	0	0.00 %	3.97%
Arg	R	0	0.00%	5.16%	Ser	S	4	7.55 %	7.12%
Thr	T	4	7.55%	5.67%	Val	V	4	7.55 %	6.58%
Trp	W	1	1.89%	1.23%	Tyr	Y	5	9.43 %	3.18%

Number of acidic (negative) amino acids (ED): 2
 3.77%
Number of basic (positive) amino acids (KR): 3
 5.66%
Total charge (KRED): 5
 9.43%
Net charge (KR - ED): 1
 1.89%
Theoretical pI: 8.18
Total linear charge density: 0.13
Average hydrophobicity: 10.81
Ratio of hydrophilicity to hydrophobicity: 0.40
Percentage of hydrophilic amino acid: 28.30%
Percentage of hydrophobic amino acid: 71.70%

Ratio of %hydrophilic to %hydrophobic:

0.39

77ORF104 sequence

```
34393      atggtaaccaaagaattttttaaaaactaaacttgagtgttcagat
1      M  V  T  K  E  F  L  K  T  K  L  E  C  S  D
34438      atgtacgctcagaaactcatagatgaggcacagggcgatgaaaat
16     M  Y  A  Q  K  L  I  D  E  A  Q  G  D  E  N
34483      aggttgtagcactatctatccaaaaacttgcagaacgtcataca
31     R  L  Y  D  L  F  I  Q  K  L  A  E  R  H  T
34528      cgccccgctatcgtcgaatattaa 34551
46     R  P  A  I  V  E  Y  *
```

Physico-chemical parameters of ORF 77ORF104

1 MVTKEFLKTK LECSDMYAQK LIDEAQGDEN RLYDLFIQKL AERHTRPAIV EY

Number of amino acids: 52
 Average molecular weight (Daltons): 6193.13
 Mean amino acid weight (Daltons): 119.10
 Monoisotopic molecular weight (Daltons): 6189.12
 Mean amino acid monoisotopic weight (Daltons): 119.02

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	4	7.69 %	7.58%	Cys	C	1	1.92%	1.66%
Asp	D	4	7.69 %	5.28%	Glu	E	6	11.54 %	6.37%
Phe	F	2	3.85 %	4.09%	Gly	G	1	1.92%	6.84%
His	H	1	1.92 %	2.24%	Ile	I	3	5.77%	5.81%
Lys	K	5	9.62 %	5.95%	Leu	L	6	11.54 %	9.42%
Met	M	2	3.85 %	2.37%	Asn	N	1	1.92%	4.45%
Pro	P	1	1.92 %	4.9%	Gln	Q	3	5.77%	3.97%
Arg	R	3	5.77 %	5.16%	Ser	S	1	1.92%	7.12%
Thr	T	3	5.77 %	5.67%	Val	V	2	3.85%	6.58%
Trp	W	0	0.00 %	1.23%	Tyr	Y	3	5.77%	3.18%

Number of acidic (negative) amino acids (ED): 10
 19.23%
 Number of basic (positive) amino acids (KR): 8
 15.38%
 Total charge (KRED): 18
 34.62%
 Net charge (KR - ED): -2
 3.85%
 Theoretical pI: 5.03
 Total linear charge density: 0.38
 Average hydrophobicity: -5.81
 Ratio of hydrophilicity to hydrophobicity: 1.47
 Percentage of hydrophilic amino acid: 53.85%
 Percentage of hydrophobic amino acid: 46.15%

164

Ratio of %hydrophilic to %hydrophobic:

1.17

77ORF182 sequence

```
29268      atgttcaatataaaaacgaaaaacggaggaagtcaagatgtattac
1      M  F  N  I  K  R  K  T  E  E  V  K  M  Y  Y
29313      gaaataggcgaaatcatacgcaaaaatattcatgttaacggattc
16     E  I  G  E  I  I  R  K  N  I  H  V  N  G  F
29358      gatTTtaagctattcattttaaaaggTcatatgggcatatcaata
31     D  F  K  L  F  I  L  K  G  H  M  G  I  S  I
29403      caagttaaagatatgaacaacgtaccaattaaacatgcttatgtc
46     Q  V  K  D  M  N  N  V  P  I  K  H  A  Y  V
29448      gtagatgagaatgacttagatatggcatcagacttatttaaccaa
61     V  D  E  N  D  L  D  M  A  S  D  L  F  N  Q
29493      gcaatagatgaatggattgaagagaacacagacgaacaggacaga
76     A  I  D  E  W  I  E  E  N  T  D  E  Q  D  R
29538      ctaattaacttagtcatgaaatggtag 29564
91     L  I  N  L  V  M  K  W  *
```

Physico-chemical parameters of ORF 77ORF182

1 MFNIKRKTEE VKMYEIGEI IRKNIHVNGF DFKLFILKGH MGISIQVKDM NNVPIKHAYV
 61 VDENDLDMAS DLFNQAIDEW IEENTDEQDR LINLVMKW

Number of amino acids: 98
 Average molecular weight (Daltons): 11691.50
 Mean amino acid weight (Daltons): 119.30
 Monoisotopic molecular weight (Daltons): 11683.84
 Mean amino acid monoisotopic weight (Daltons): 119.22

Amino acid composition

Aci d	Symbo l	Numb er	%	Average % in Swissprot	Aci d	Symbo l	Numb er	%	Average % in Swissprot
Ala	A	3	3.06%	7.58%	Cys	C	0	0.00%	1.66%
Asp	D	9	9.18%	5.28%	Glu	E	9	9.18%	6.37%
Phe	F	5	5.10%	4.09%	Gly	G	4	4.08%	6.84%
His	H	3	3.06%	2.24%	Ile	I	12	12.24%	5.81%
Lys	K	9	9.18%	5.95%	Leu	L	6	6.12%	9.42%
Met	M	6	6.12%	2.37%	Asn	N	9	9.18%	4.45%
Pro	P	1	1.02%	4.9%	Gln	Q	3	3.06%	3.97%
Arg	R	3	3.06%	5.16%	Ser	S	2	2.04%	7.12%
Thr	T	2	2.04%	5.67%	Val	V	7	7.14%	6.58%
Trp	W	2	2.04%	1.23%	Tyr	Y	3	3.06%	3.18%

Number of acidic (negative) amino acids (ED): 18
 18.37%
 Number of basic (positive) amino acids (KR): 12
 12.24%
 Total charge (KRED): 30
 30.61%
 Net charge (KR - ED): -6
 6.12%
 Theoretical pI: 4.76
 Total linear charge density: 0.33
 Average hydrophobicity: -3.89
 Ratio of hydrophilicity to hydrophobicity: 1.28

167

Percentage of hydrophilic amino acid:	51.02%
Percentage of hydrophobic amino acid:	48.98%
Ratio of %hydrophilic to %hydrophobic:	1.04

Table 5

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100017|lan|77ORF017 Phage 77 ORF |23269-23982|-3
(237 letters)

Database: nr
393,678 sequences; 120,452,765 total letters

		Score (bits)	E Value
Sequences producing significant alignments:			
gi 4493986 emb CAB39045.1	(AL034559) predicted using hexExon; ...	41	0.010
gi 730607 sp P23250 RPI1_YEAST	NEGATIVE RAS PROTEIN REGULATOR P...	38	0.053
gi 3097044 emb CAA75299	(Y15035) K1R [Cowpox virus]	38	0.090
gi 2146245 pir S73794	hypothetical protein H91_orf180 - Mycopl...	38	0.090
gi 83910 pir S04682	ribosomal protein var1 - yeast (Candida gl...	37	0.15
gi 133135 sp P21358 RMAR_CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN ...	37	0.15
gi 2128843 pir H64475	hypothetical protein MJ1409 - Methanococ...	36	0.20
gi 5107017 gb AAD39926.1 AF126285_2	(AF126285) RNA polymerase [...	36	0.35
gi 2146210 pir S73342	hypothetical protein E07_orf166 - Mycopl...	35	0.60

Database: swissprot
79,449 sequences; 28,874,452 total letters

		Score (bits)	E Value
Sequences producing significant alignments:			
sp P23250 RPI1_YEAST	NEGATIVE RAS PROTEIN REGULATOR PROTEIN.	38	0.014
sp P21358 RMAR_CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	37	0.040
sp Q21444 LDLC_CAEEL	LDLC PROTEIN HOMOLOG.	34	0.35
sp P27240 RFAY_ECOLI	LIPOPOLYSACCHARIDE CORE BIOSYNTHESIS PROT.	33	0.46
sp P53192 YGCO_YEAST	HYPOTHETICAL 27.1 KD PROTEIN IN ALK1-CKB1.	33	0.60
sp P32908 SMC1_YEAST	CHROMOSOME SEGREGATION PROTEIN SMC1 (DA-B.	33	0.60
sp P54683 TAGB_DICDI	PRESTALK-SPECIFIC PROTEIN TAGB PRECURSOR .	32	0.78
sp Q03100 CYAA_DICDI	ADENYLATE CYCLASE, AGGREGATION SPECIFIC (.	32	0.78

169

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100019|lan|77ORF019 Phage 77 ORF|39851-40501|2
(216 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

		Score	E
		(bits)	Value
Sequences producing significant alignments:			
gi 3341966 dbj BAA31932	(AB009866) orf 59 [bacteriophage phi PVL]	437	e-122
gi 2689911	(AE000792) B. burgdorferi predicted coding region BB...	38	0.058
gi 1171589 emb CAA64574	(X95275) frameshift [Plasmodium falcip...	37	0.10
gi 4493986 emb CAB39045.1	(AL034559) predicted using hexExon; ...	36	0.23
gi 141257 sp P18019 YPI9_CLOPE	HYPOTHETICAL 14.5 KD PROTEIN (OR...	36	0.29
gi 133412 sp P27059 RPOB_ASTLO	DNA-DIRECTED RNA POLYMERASE BETA...	35	0.51
gi 3122231 sp Q58851 HISX_METJA	HISTIDINOL DEHYDROGENASE (HDH) ...	35	0.51
gi 3649757 emb CAB11106.1	(Z98547) predicted using hexExon; MA...	34	0.66
gi 2688313	(AE001146) sensory transduction histidine kinase, pu...	34	0.87

Database: swissprot
79,449 sequences; 28,874,452 total letters

		Score	E
		(bits)	Value
Sequences producing significant alignments:			
sp P18019 YPI9_CLOPE	HYPOTHETICAL 14.5 KD PROTEIN (ORF9).	36	0.079
sp Q58851 HISX_METJA	HISTIDINOL DEHYDROGENASE (EC 1.1.1.23) (H.	35	0.14
sp P27059 RPOB_ASTLO	DNA-DIRECTED RNA POLYMERASE BETA CHAIN (E.	35	0.14
sp Q02224 CENE_HUMAN	CENTROMERIC PROTEIN E (CENP-E PROTEIN).	34	0.31
sp P04931 ARP_PLAFA	ASPARAGINE-RICH PROTEIN (AG319) (ARP) (FRA..	33	0.53
sp P18011 IPAB_SHIFL	62 KD MEMBRANE ANTIGEN.	32	0.69
sp P18709 VTA2_XENLA	VITELLOGENIN A2 PRECURSOR (VTG A2) [CONTA..	32	0.90
sp Q64409 CP3H_CAVPO	CYTOCHROME P450 3A17 (EC 1.14.14.1) (CYPI..	32	0.90
sp P21358 RMAR_CANGA	MITOCHONDRIAL RIBOSOMAL PROTEIN VAR1.	32	0.90
sp Q03945 IPAB_SHIDY	62 KD MEMBRANE ANTIGEN.	32	1.2

170

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100043|lan|77ORF043 Phage 77 ORF|29304-29564|3
(86 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 3341947 dbj BAA31913 (AB009866) orf 39 [bacteriophage phi PVL]	182	6e-46
gi 744518 prf 2014422A FKBP-rapamycin-associated protein [Homo...]	32	0.84
gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN...	32	0.84
gi 1169735 sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTE...	32	0.84
gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa...]	32	0.84
gi 3875402 emb CAA98122 (Z73906) cDNA EST EMBL:D64544 comes fr...	31	2.5
gi 1084792 pir S54091 hypothetical protein YPR070w - yeast (Sa...]	30	4.2

Database: swissprot
79,449 sequences; 28,874,452 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) .	32	0.24
sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R.	32	0.24
sp P34554 YNP1 CAEEL HYPOTHETICAL 42.2 KD PROTEIN T05G5.1 IN C.	28	3.5
sp Q24118 LIO_DROME LINOTTE PROTEIN.	28	3.5
sp P80034 ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	28	3.5
sp P22922 A1AT_BOMMO ANTITRYPSIN PRECURSOR (AT).	28	3.5
sp Q44363 TRAA_AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	28	3.5
sp P38255 YBU5_YEAST HYPOTHETICAL 51.3 KD PROTEIN IN PHO5-VPS1.	27	6.0
sp P55822 SH3B_HUMAN SH3BGR PROTEIN (21-GLUTAMIC ACID-RICH PRO.	27	7.9
sp Q58482 YA82_METJA HYPOTHETICAL PROTEIN MJ1082.	27	7.9
sp P34252 YKK8_YEAST HYPOTHETICAL 52.3 KD PROTEIN IN HAP4-AAT1.	27	7.9

171

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100102|lan|77ORF102 Phage 77 ORF|29051-29212|2
(53 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

	Score (bits)	E Value
Sequences producing significant alignments:		
gi 3341946 dbj BAA31912 (AB009866) orf 38 [bacteriophage phi PVL]	96	3e-20
gi 4325288 gb AAD17315 (AF123593) voltage-dependent sodium cha...	28	7.1
gi 2649684 (AE001040) A. fulgidus predicted coding region AF092...	28	9.3

Database: swissprot
79,449 sequences; 28,874,452 total letters

	Score (bits)	E Value
Sequences producing significant alignments:		
sp P42087 HUTM_BACSU PUTATIVE HISTIDINE PERMEASE.	26	7.1
sp P04775 CIN2_RAT SODIUM CHANNEL PROTEIN, BRAIN II ALPHA SUBU...	26	9.2
sp P42619 YQJF_ECOLI HYPOTHETICAL 17.2 KD PROTEIN IN EXUR-TDCC...	26	9.2

172

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|100104|lan|77ORF104 Phage 77 ORF|34393-34551|1
(52 letters)

Database: nr
373,355 sequences; 114,214,446 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 2315523 (AF016452) similar to the leucine-rich domains found...	29	4.2
gi 4377168 gb AAD18990 (AE001666) CT711 hypothetical protein [...	29	5.4
gi 3882171 dbj BAA34445 (AB018268) KIAA0725 protein [Homo sapi...	28	9.3

Database: swissprot
79,449 sequences; 28,874,452 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P04879 RRPP_VSVIG RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.	27	5.4
sp P04880 RRPP_VSVIM RNA POLYMERASE ALPHA SUBUNIT (EC 2.7.7.48.	27	5.4
sp Q13946 CN7A_HUMAN HIGH-AFFINITY CAMP-SPECIFIC 3',5'-CYCLIC .	26	7.1
sp P35381 ATPA_DROME ATP SYNTHASE ALPHA CHAIN, MITOCHONDRIAL P.	26	9.3
sp P54659 MVPB_DICDI MAJOR VAULT PROTEIN BETA (MVP-BETA).	26	9.3
sp P40397 YHXC_BACSU HYPOTHETICAL OXIDOREDUCTASE IN APRE-COMK .	26	9.3

173

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|122748|lan|77ORF182 Phage 77 ORF|29268-29564|3
(98 letters)

Database: nr
393,678 sequences; 120,452,765 total letters

Sequences producing significant alignments:	Score (bits)	E Value
gi 3341947 dbj BAA31913.1 (AB009866) orf 39 [bacteriophage phi..	182	8e-46
gi 1084792 pir S54091 hypothetical protein YPR070w - yeast (Sa..	35	0.13
gi 1169736 sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN..	32	1.1
gi 744518 prf 2014422A FKBP-rapamycin-associated protein [Homo..	32	1.1
gi 5051381 emb CAB44736.1 (AL049653) dJ647M16.2 (FK506 binding..	32	1.1
gi 4826730 ref NP_004949.1 pFRAP1 FK506 binding protein 12-rap..	32	1.1
gi 3282239 (U88966) rapamycin associated protein FRAP2 [Homo sa..	32	1.1

Database: swissprot
79,909 sequences; 29,054,478 total letters

Sequences producing significant alignments:	Score (bits)	E Value
sp P42345 FRAP_HUMAN FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) .	32	0.29
sp P42346 FRAP_RAT FKBP-RAPAMYCIN ASSOCIATED PROTEIN (FRAP) (R.	32	0.29
sp P40557 YIA5_YEAST PUTATIVE DISULFIDE ISOMERASE YIL005W PREC.	29	3.3
sp Q24118 LIO_DROME LINOTTE PROTEIN.	28	4.4
sp Q44363 TRAA_AGRT6 CONJUGAL TRANSFER PROTEIN TRAA.	28	4.4
sp P80034 ACH2_BOMMO ANTICHYMOTRYPSIN II (ACHY-II).	28	4.4
sp P34554 YNP1_CAEEL HYPOTHETICAL 42.2 KD PROTEIN T05G5.1 IN C.	28	4.4
sp P22922 A1AT_BOMMO ANTITRYPSIN PRECURSOR (AT).	28	4.4

Table 6

1st position (5' end)	2nd position				3rd position (3' end)
	U	C	A	G	
U	Phe	Ser	Tyr	Cys	U
	Phe	Ser	Tyr	Cys	C
	Leu	Ser	Stop	Stop	A
	Leu	Ser	Stop	Trp	G
C	Leu	Pro	His	Arg	U
	Leu	Pro	His	Arg	C
	Leu	Pro	Gln	Arg	A
	Leu	Pro	Gln	Arg	G
A	Ile	Thr	Asn	Ser	U
	Ile	Thr	Asn	Ser	C
	Ile	Thr	Lys	Arg	A
	Met	Thr	Lys	Arg	G
G	Val	Ala	Asp	Gly	U
	Val	Ala	Asp	Gly	C
	Val	Ala	Glu	Gly	A
	Val	Ala	Glu	Gly	G

Table 7

Bacteriophage 3A, complete genome sequence

1	caaacgctag	caacgcggat	aaatttttca	tgaaggggg	tcttttatg	aagttaacaa	aaaaacagct
71	aaaagaatat	atagaagatt	acaaaaaatc	tgatgacata	ttaatttaatt	tgtatataga	aacatatgaa
141	ttttattgtc	ggttaagaga	tgaacttaaa	aatagtgtat	taatgataga	gcatacaaac	aaggctgggtg
211	cgagcaatat	tattaagaat	ccattaaagca	tagaactgac	aaaaacagtt	caaacactaa	ataacttact
281	caagtctatg	ggtttaactg	cagcacaag	aaaaaagata	gttcaagaag	aagggtggatt	cggtgactat
351	taaagtttta	aatgaacctt	cacaaaaact	attaacaaca	tggtatgcag	agcaagtcac	tcaagggaaa
421	ataaaaacaa	gcaaatatgt	tagaaaaaaa	tgtgagagac	atcttagata	tctagaaaaat	ggaggttaaat
491	gggtatttga	tgaagaatta	gcgcatcgct	ctattcgatt	tatagaaaaa	ttttgttaaac	cttccaaagg
561	atctaaacgt	caacttgtat	tacagccatg	gcaacatttt	attatcgcca	gtttgtttgg	ttgggttcat
631	aaagaaacaa	aactgcgag	gtttaaagaa	gctttgatatt	tatatgggag	aaaaaatggt	aaaaacaacca
701	ctattttctg	gggttgcatac	tatgctgtat	cacaagatgg	agaaaaatggt	gcagaaattc	atttgttagc
771	aaacgttaag	aaacaagcta	ggatttctatt	tgtatgaact	aaggcgatga	ttaaagctag	cccaagctt
841	gataaaaaatt	tcagaacatt	aagagatgaa	atccattatg	acgcaacgat	atcaaaaaatt	atgccccaaag
911	catcagatag	cgataagtta	gatggattga	atacacacat	ggggattttt	gatgaatttc	atgaatttaa
981	agactataaa	ttgatttcag	ttataaaaaa	ctcaagagct	gcaagggttac	aacctcttct	catctacatt
1051	acgacagcag	gggtatcaatt	agatgggtcca	cttgttgata	tggtagaagc	gggaagagac	accttagatc
1121	aaatcataga	agacgaaaga	actttttatt	atcttagcatc	tttggtatgat	gacgatgata	ttaatgatcc
1191	gtcgaaactg	ataaaagcaa	atcccaactt	aggtgtctct	ataaatttag	atgagatgaa	agaagagtgg
1261	gaaaaagcta	agagaacacc	agctgaacgt	ggagatttta	taaccaaaaag	gtttaaatatc	tttgcttaata
1331	atgacgagat	gagttttatt	gattacccaa	cactccaaaa	aaataatgaa	attgtttctt	tagaagagct
1401	ggaaggcaga	ccgtgcacga	ttggttatga	tttatcagaa	acagaggact	ttacagccgc	gtgtgctact
1471	tttgcgttag	ataatggtaa	agttgcagtt	ttatcgcatc	catggattcc	taagcacaaa	gttgaattat
1541	ctaacgaaaa	aataccctat	agagaatggg	aagaagatgg	cttattaaca	gtgcaagata	agccttatat
1611	tgactaccaa	gatgttttaa	attggataat	taagatgaat	gagcattatg	tagtagaaaa	aattacttat
1681	gatagagcga	acgcattcaa	actaaatcaa	gagttaaaaa	attacgggtt	tgaacgggaa	gaaacaagac
1751	aaggagcttt	gaccttgagc	ccgtgcattga	aggattttaa	agaaatgttt	ttagatggga	aaataatatt
1821	taataataat	cctttaatga	aatggatat	caataatgtt	cagttgaaac	tagacagaaa	cggaaactgg
1891	ttgccgtcta	agcaaaagcag	atatacgtaa	atagatggct	ttgcagcatt	tttaaacaca	tatacagata
1961	ttatgaataa	agttgtttct	gatagtgggt	aaggaaacat	agagtttatt	agtattaaag	acataatgcg
2031	ttaggagggt	gaatgtttatc	gcaaaagaga	atattgtcac	acgcataaag	aaaaaattga	tagacaattg
2101	gattgatcag	tcacttctta	agctttatga	ctttagccca	tggaaaaaata	gatctttttg	gggtgttaatt
2171	aataatacgc	ttgaaactaa	tgaacagata	ttttcagcta	ttacaaaagt	atctaattcg	atggctagtt
2241	tgcccttgaa	aatgtatgaa	gattataaag	tagttaatag	agaagtatct	gatttactta	cagtgctacc
2311	gaataaattct	ctgagcagtt	ttgattttat	taatcaaat	gaaacaatca	gaaatgaaaa	aggttaattgca
2381	tatgtgctaa	ttgaacgaga	catctatcat	caaccatcaa	agcttttctt	atataaatcca	gatgttggtg
2451	aaatgttaatt	tgaaaaccaa	tcacgtgaac	ttttattatc	catctatgct	gcaactggaa	ataaattgat
2521	tggtcataat	atggacatgt	tgcattttaa	acacatcgtg	gcattctaata	tggtgcaagg	cattagtccg
2591	attgatgtgt	tgaagaatac	aaactgatttt	gataatgcag	taagaacctt	taactctaca	gaaatgcaaa
2661	aacctgattc	tttctgctct	aaatatgggt	ccaatgtagg	taaaagaaaa	agggcagcaag	tgttagaaga
2731	tttcaaacag	tactatgaag	aaaacgggtg	aatattatct	caagagcctg	gtgttgaaat	cgaaccgtta
2801	cctaaaaaat	atgtctctga	agatatagtg	gcaagcgaga	atttaacaag	agaaagagta	gctaaccgttt
2871	ttcaattgcc	ctcagtatct	ttaaatgcaa	gatcaaatca	aaatttcgag	aaaaatgaag	agttaaacag
2941	attttactgt	cagcatacct	tattgccaat	cgtcaaacag	tatgaagaag	aatttaactcg	gaaactactt
3011	actaaacag	acagagaaaa	aaataggtat	tttaaattta	acgttaaatc	ttatttaagg	gctgatagtg
3081	caacacaagc	agaaggtgac	tttaaagcag	ttcgtagtgg	ttactacact	ataaatgaca	ttagagagtg
3151	ggaagattta	caaccagttg	aaggtggaga	taagccgcta	ataagcggtg	atttataccc	aattgacacg
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3291	aaaagaaaaat	caaaaagtaa	aggtgaaata	ttttattatg	gtgatattgt	aagtataaaa	tggtttgaaa
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Table 8

Bacteriophage 3A ORFs list

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100382	3AORF004	3	17457..19370	637	gctattttattagaaaggaaggtgc	att	taa
100383	3AORF005	1	334..2034	566	agaaaaagatagttcaagaagaag	gtg	taa
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100727	3AORF349	-2	6882..6983	33	tagaacgaccaataactgtatttag	atc	taa
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189

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Table 9

Bacteriophage 96, complete genome sequence

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Table 10

Bacteriophage 96 ORFs list

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100738	96ORF006	2	40589..42043	484	aatgatttagggtaggtgttgacca	atg	tag
100739	96ORF007	1	18652..20091	479	tatacacacataactaaacctgaacg	att	tga
100740	96ORF008	2	8960..10201	413	tggcagaatttggggcgataacga	atg	tga
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100954	96ORF222	-1	39860..39997	45	acttttatttaggttcaactccatt	att	taa
100955	96ORF223	-1	24716..24853	45	acatttcaaatgattctggaacaac	ata	taa
100956	96ORF224	-2	26794..26931	45	caatatcacgccatgtagtttttaa	ctg	taa
100957	96ORF225	-2	19201..19338	45	caaacaatggattgtaatacaataa	atg	tga
100958	96ORF226	-2	15709..15846	45	tgacttgcgtgtgtgtctaacacaat	ata	taa
100959	96ORF227	-3	36711..36848	45	acattgactgccccgataattatct	ata	tga
100960	96ORF228	3	2325..2459	44	tcgccatagtgagttccaataccgt	ata	taa
100961	96ORF229	-1	38612..38746	44	ttgtcattgataacctattcttatag	atg	tga
100962	96ORF230	-1	31733..31867	44	gctggattgtagtggttaagtaaat	ctg	tag
100963	96ORF231	-2	12076..12210	44	tgactcatagcttcaactgttctgt	ctg	taa
100964	96ORF232	-3	31644..31778	44	atagtcctcaagtgttaacctagt	ttg	taa
100965	96ORF233	-3	23988..24122	44	atttgattgtgaagttcaggctcaa	ctg	taa
100966	96ORF234	-3	17529..17663	44	agtagcgttttttgaaatcgtaacct	atg	taa
100967	96ORF235	1	7153..7284	43	aatgctaaggtccaatagaaatca	atg	tag
100968	96ORF236	2	2681..2812	43	ttctttcacttcaacttcacatttc	ata	tga
100969	96ORF237	2	4496..4627	43	gtactatgcttcacagctcttagcga	ttg	taa
100970	96ORF238	-1	41720..41851	43	cacctgtaattcttgaaatagttga	ata	tga
100971	96ORF239	-1	35324..35455	43	acttactaataaaatagaatagttt	gtg	taa
100972	96ORF240	-1	8570..8701	43	atccccgttttgacttaatacatca	atc	tga
100973	96ORF241	-2	33502..33633	43	ataattttgtaataactcttagggat	atg	tag
100974	96ORF242	-2	23662..23793	43	agctaagtctacagcagtggtgtaa	atc	tag
100975	96ORF243	-3	32391..32522	43	acctggagcagcttgcgtcatataa	ata	tag
100976	96ORF244	-3	30273..30404	43	aaaacttctgtataactcttggtaa	atc	tga
100977	96ORF245	-3	5895..6026	43	tgactaataatgcttataattctta	atc	taa
100978	96ORF246	-3	2679..2810	43	attcatcaagaaactatagccggtc	atg	tga
100979	96ORF247	1	34891..35019	42	acatcaagcaaatctggtgtgttag	ttg	taa
100980	96ORF248	2	30668..30796	42	aattattacattaaagctggtgtga	atg	tag
100981	96ORF249	2	31838..31966	42	caaatattagcttgtagtgagttag	atg	taa
100982	96ORF250	2	33539..33667	42	cttaccagaaacagcacaggtagaa	ata	taa
100983	96ORF251	-1	20486..20614	42	cttctgtacgagccacacgcaatga	ttg	tag
100984	96ORF252	-1	15128..15256	42	gataatttcattactagctactacta	ata	tga
100985	96ORF253	-2	41446..41574	42	aaaacctaatcagataaacgataa	ttg	tga
100986	96ORF254	-2	41005..41133	42	gttaraaccatgaccggctacaagc	ata	taa
100987	96ORF255	-2	23008..23136	42	aggataaatgacttgaccatcttc	ata	taa
100988	96ORF256	-2	14794..14922	42	ttgtatgcgtcaatgagttggtcga	ttg	tag
100989	96ORF257	-2	8503..8631	42	tacctaaccttttttaataatttcta	atg	tga
100990	96ORF258	-3	22143..22271	42	aaacgctttgtaaaatgcctctgca	att	tga
100991	96ORF259	-3	18639..18767	42	cttgratctattatagagattaacc	att	tag
100992	96ORF260	-3	15624..15752	42	gttttggtaactagccactgtatag	ata	taa
100993	96ORF261	2	18746..18871	41	catattgaggctctaatagagtcac	ata	taa
100994	96ORF262	-1	13067..13192	41	aattaattaattcttctctgtgtgg	ttg	taa
100995	96ORF263	-2	18742..18867	41	taacagacacgtctaatcgccctac	att	tga
100996	96ORF264	-2	18376..18501	41	catattatcataaagaacaagtaac	ttg	taa
100997	96ORF265	-2	367..492	41	ctaaacgaaaaagagggtacaatac	atc	tga
100998	96ORF266	-3	32802..32927	41	aggatatatccatttgatacaatact	ttg	taa
100999	96ORF267	-3	10194..10319	41	atcatcgaaaggcgataactcgtaa	ttg	tga
101000	96ORF268	1	1159..1281	40	ttattcttctcttttgtaattgtaa	atg	taa
101001	96ORF269	2	10373..10495	40	gacagagttgaaaaagaaatcatga	atg	taa
101002	96ORF270	2	15734..15856	40	ttattcggcgtaatcgactgatgc	ttg	tag
101003	96ORF271	-1	43451..43573	40	c c tNo shine-dalgarno sequence	att	tga
101004	96ORF272	-1	36959..37081	40	acgctataaaaaataactttttattag	atg	tag
101005	96ORF273	-1	35798..35920	40	ctgacgcactttgtgtggtttgtagc	att	taa
101006	96ORF274	-1	8147..8269	40	ctgctctctctatgtttgttagtct	ctg	tga
101007	96ORF275	-2	43066..43188	40	tttaacttactaactttcttttgat	ata	tga
101008	96ORF276	-2	42535..42657	40	aaataatgtaaatgttttcatagt	att	tag
101009	96ORF277	-2	30628..30750	40	tttgtagtcccgctctcgcaaaagt	ctg	taa

101010	96ORF278	-2	13291..13413	40	ttcgtatcttccaagcaattcattt	ttg	tga
101011	96ORF279	-2	3172..3294	40	cagattgttttagtaacgcctaattt	atc	taa
101012	96ORF280	-3	18804..18926	40	taataataaccaacacgtgtatcaaca	att	tag
101013	96ORF281	-3	15843..15965	40	atttaaaaagtgtattctataacca	atc	tag
101014	96ORF282	-3	8460..8582	40	ttagtcatcactcaattctttttcc	att	taa
101015	96ORF283	-3	7593..7715	40	gatgttgtctacacagtgctaacac	atg	taa
101016	96ORF284	-3	6453..6575	40	aattaatttttaattaccatttcta	att	tga
101017	96ORF285	1	15082..15201	39	caatacttagtcacaacattcaag	att	taa
101018	96ORF286	1	34444..34563	39	acacaaacgttaataagcaaaagtga	atg	tag
101019	96ORF287	2	27920..28039	39	cctattttagcagttgttgcagtaa	ttg	tag
101020	96ORF288	2	28415..28534	39	atcggtcttttaactggcgtaataga	atc	tag
101021	96ORF289	2	38147..38266	39	tatcaaatgcttaatttaggcaagt	atc	tga
101022	96ORF290	3	40917..41036	39	gcaaatttaaacactttcacatcat	atg	taa
101023	96ORF291	-2	38815..38934	39	tctctaaaaacagcttacagcgaac	ata	taa
101024	96ORF292	-2	32671..32790	39	ctataggattataaatcgctgacgt	ata	tga
101025	96ORF293	-2	31216..31335	39	ttgatttgatgtttcttatactga	ttg	taa
101026	96ORF294	-2	21589..21708	39	gtatcttcacagaaatcgcttaaaa	atc	taa
101027	96ORF295	-2	18976..19095	39	tatcaatatatgctaaccctagcacc	ata	taa
101028	96ORF296	-2	11482..11601	39	gccacctcgactctttttgcaacc	att	taa
101029	96ORF297	-3	12933..13052	39	tcacgaaataatgtttctttaattt	ata	taa
101030	96ORF298	-3	8262..8381	39	gaactgatcttgcttaaatgattta	att	tag
101031	96ORF299	-3	6993..7112	39	cattagcattagcgaatgggttga	ttg	tga
101032	96ORF300	2	23516..23632	38	actacatctgaacaactaaaatttc	atc	tag
101033	96ORF301	2	25943..26059	38	agattagaagaagaaaaagaagac	gtg	taa
101034	96ORF302	2	36929..37045	38	tattgggggtttgttaacatggggca	atg	tag
101035	96ORF303	3	4476..4592	38	ataaaagctacctagtagcagta	atg	tga
101036	96ORF304	3	20586..20702	38	tactctaagatagctaaagcaatac	gtg	tga
101037	96ORF305	3	28356..28472	38	cggttaccaatgtgcttgatacgat	ttg	taa
101038	96ORF306	-1	24359..24475	38	acttaataaaaagccgtatcgtgcc	atg	taa
101039	96ORF307	-1	20147..20263	38	ttgtacctatacagagtaactcctt	att	tag
101040	96ORF308	-2	38158..38274	38	ttccgtatccactttctaagaagc	gtg	tga
101041	96ORF309	-2	35149..35265	38	agcttgtttgtatcgtcttcaacga	ata	taa
101042	96ORF310	-2	31423..31539	38	gtaatatgattaggtctcctcttat	ttg	taa
101043	96ORF311	-2	10438..10554	38	cgcttttaaatcgtttaggtcact	atc	taa
101044	96ORF312	-2	1390..1506	38	gagaacaacacacaacattacaaca	atc	taa
101045	96ORF313	-3	33051..33167	38	acgtcctggtttctagatcgtaatac	ata	tag
101046	96ORF314	-3	25194..25310	38	agcaaacctgtaaaagataacattga	atc	taa
101047	96ORF315	-3	6273..6389	38	cattcttgcctaacacgtcagattga	ctg	tga
101048	96ORF316	-3	4281..4397	38	ataattcgtattcattaatcattaa	att	tag
101049	96ORF317	1	2260..2373	37	atgactccttttctcatatttcttt	ata	taa
101050	96ORF318	2	21230..21343	37	atttcacactttttagttgtctct	ttg	taa
101051	96ORF319	3	18018..18131	37	atactgagtcaccaatttaagctcg	atg	tag
101052	96ORF320	3	36972..37085	37	attacagatatcctaaggggttccg	att	taa
101053	96ORF321	-1	36302..36415	37	ctcttgagttttttgacctaattta	atc	taa
101054	96ORF322	-1	32606..32719	37	ccataagttattttctccagttctat	att	taa
101055	96ORF323	-1	11453..11566	37	ttaaaccgttcttttttatcaattc	att	tga
101056	96ORF324	-1	7268..7381	37	tactgggttcgcccagtggaagttct	ata	tga
101057	96ORF325	-2	32347..32460	37	ttactgcattttgtatatggcgataa	atc	tag
101058	96ORF326	-2	24682..24795	37	acgtttattacgctcataaagccat	ata	tag
101059	96ORF327	-2	23905..24018	37	aaatggctgtggcgcttgaccatat	gtg	taa
101060	96ORF328	-2	21460..21573	37	agagcactaatacgtttttgttctt	ctg	tga
101061	96ORF329	-2	21208..21321	37	gacttaacttcttctgatattcata	atc	tga
101062	96ORF330	-2	18085..18198	37	ccagtcgacaccagcaagattattct	ttg	tag
101063	96ORF331	-2	8170..8283	37	actttgagacgtcgtctgtctctct	atg	tag
101064	96ORF332	-2	5971..6084	37	caatttggtttccggttttctcttag	ttg	tag
101065	96ORF333	-3	37632..37745	37	acettgcttaatacaagtcgtaatta	att	tga
101066	96ORF334	-3	29628..29741	37	ctgagttagtggttgtaaaatgtcat	ttg	tag
101067	96ORF335	-3	7164..7277	37	ttageggatatccgttttctagtaa	atc	taa
101068	96ORF336	1	22903..23013	36	gtaaaaaaagacaatatgactatta	ctg	tga
101069	96ORF337	1	43258..43368	36	taattgacgtgggtatttttttaggt	ttg	taa
101070	96ORF338	2	12668..12778	36	gaactgggtggaatggcgatggaaca	atc	tag
101071	96ORF339	2	28292..28402	36	ttcactgcttttaattcagttgctta	ctg	taa
101072	96ORF340	2	35396..35506	36	ttcctaataagaacataagtcaacggt	att	tga
101073	96ORF341	3	25428..25538	36	actcgagaacaattagaaaaagcaa	ttg	tga
101074	96ORF342	-1	40913..41023	36	tatctgggaaatttaacttaataaa	ata	tga
101075	96ORF343	-1	39173..39283	36	tgccacatttttagtgtcaggattga	ttg	taa
101076	96ORF344	-1	37580..37690	36	gggtctaccttttaacgctcgtttcag	ata	taa
101077	96ORF345	-1	31556..31666	36	ggattattcttttctaataacttcaa	ttg	tga
101078	96ORF346	-1	29972..30082	36	ggctactccttatcctaataataaat	ttg	taa
101079	96ORF347	-1	28787..28897	36	ctgccaaagctctgtagcaattactt	ttg	tga
101080	96ORF348	-1	21839..21949	36	ttaaaatccgataaaaataacattgc	ctg	tga
101081	96ORF349	-1	3647..3757	36	taaaacttcgaagttaccagcgt	ttg	tga

101082	96ORF350	-2	40801..40911	36	accattccaattttgcccataatgat	gtg	tag
101083	96ORF351	-2	38953..39063	36	tatcttttaaaattctcgtaatagc	atc	taa
101084	96ORF352	-2	31585..31695	36	tagctgtcatcactagtatttttga	atc	taa
101085	96ORF353	-2	24550..24660	36	atagtcggttttaccgcctcgtact	att	tag
101086	96ORF354	-2	20083..20193	36	atcatcatttttgatatttctcaaac	ata	tga
101087	96ORF355	-2	991..1101	36	gcattctggcagtagcagcgtaaaac	atc	tag
101088	96ORF356	-3	38148..38258	36	taagaaaagcgtgcgcgatacaataa	att	tga
101089	96ORF357	-3	8790..8900	36	tgaagttatctagcgtatttttct	ttg	tag
101090	96ORF358	-3	4458..4568	36	ttcataaaagtattctttgtagtat	atg	tag
101091	96ORF359	1	4666..4773	35	ttatcaaaatatacaacttaattaa	atc	tag
101092	96ORF360	1	11569..11676	35	ataaatttaccgaacatgaaaatga	att	tga
101093	96ORF361	2	6122..6229	35	ggaaaacaaattgatgttgtagta	ttg	taa
101094	96ORF362	-1	40418..40525	35	ttcgtaggtgtcattacttctttaa	ttg	tag
101095	96ORF363	-1	34358..34465	35	gttttgcttgatttcgatttggtga	atg	tga
101096	96ORF364	-1	20654..20761	35	ctatttccactgattccccatctaa	atg	tga
101097	96ORF365	-1	8423..8530	35	tcttttttagagttacgaggtttca	att	tag
101098	96ORF366	-1	2402..2509	35	tgacgtatggcaacatttttagatca	atc	taa
101099	96ORF367	-2	36607..36714	35	aaaataaaaaagccagtgcgaagca	ctg	tag
101100	96ORF368	-2	27061..27168	35	caaatcgctctgcagcgttcaataa	atc	tag
101101	96ORF369	-2	26470..26577	35	atgagttgttaagttaccaccaaat	atc	taa
101102	96ORF370	-2	10327..10434	35	ccgtgccatcttctcggtataagta	ata	taa
101103	96ORF371	-2	8650..8757	35	gggtacgggtgttactgttgatat	atc	taa
101104	96ORF372	-3	14382..14489	35	gttcttttaattgatctactgttaa	att	taa
101105	96ORF373	-3	8151..8258	35	atgtttgttagtctctgtgtagtct	atg	taa
101106	96ORF374	-3	5007..5114	35	aaacgatttaagtggacattattc	ata	taa
101107	96ORF375	2	30563..30667	34	cgattagaaatctttaaanaaggac	ttg	tga
101108	96ORF376	-1	19916..20020	34	tctatgtcaggttaattgtcattaa	att	taa
101109	96ORF377	-1	9236..9340	34	ctttctgttagtaattgttttaa	atc	taa
101110	96ORF378	-1	9026..9130	34	actctttatcttttagttgcttttaa	ata	tag
101111	96ORF379	-2	28447..28551	34	cttttgtgataataaagttagtgct	ttg	tga
101112	96ORF380	-3	40329..40433	34	ccatttaccttcttgagatgttga	ttg	tga
101113	96ORF381	-3	39801..39905	34	caaaagatgaaggctttttccatc	ttg	taa
101114	96ORF382	-3	33831..33935	34	atgttggtttgtaactcgattaaagt	atc	tga
101115	96ORF383	-3	33687..33791	34	gttattacgtcttaatacttgtgtt	gtg	tag
101116	96ORF384	-3	13530..13634	34	tatacgcactagtagtgcactga	ttg	taa
101117	96ORF385	-3	3843..3947	34	tttgattgattgttctagttaagaa	att	taa
101118	96ORF386	1	12256..12357	33	agtcataaagaagtttagcaatgtga	ttg	tag
101119	96ORF387	2	2207..2308	33	tccaagactctttaactgttaactt	atc	tag
101120	96ORF388	2	2519..2620	33	attgttgaatttcgattgatctaaa	atg	tga
101121	96ORF389	2	22517..22618	33	agaagtaaaatgcgtaatgcttag	atg	tag
101122	96ORF390	2	27302..27403	33	ttccaaaattgggctaatagtgtag	ctg	taa
101123	96ORF391	2	32384..32485	33	actaaaagggttgagaaagctgtag	atg	taa
101124	96ORF392	2	39287..39388	33	aaaaacgggtactgtagatcaatca	atc	tag
101125	96ORF393	3	18153..18254	33	gtagtatatgccgactttgatttga	atg	taa
101126	96ORF394	3	24189..24290	33	tcagaccctaacattaaacaaactag	ttg	tga
101127	96ORF395	-1	15266..15367	33	tcgataaatttgtagcttgtttta	atg	tag
101128	96ORF396	-2	32239..32340	33	ttttagtgaagcatctagtgttga	ata	tag
101129	96ORF397	-2	16123..16224	33	ttatgtgtgcctatcatattaacaa	ttg	tag
101130	96ORF398	-2	13648..13749	33	tctttaactgaatgttgaatagcat	ttg	tag
101131	96ORF399	-2	10987..11088	33	acttctgtaggtattcttatatcaa	ttg	tga
101132	96ORF400	-2	3382..3483	33	cttactggtaattcttcaaaattaa	atg	taa
101133	96ORF401	-3	40794..40895	33	ccatatgatgtgaaagtgtttaa	ttg	taa
101134	96ORF402	-3	39978..40079	33	atattcctaatacacttgaacctaa	att	tga
101135	96ORF403	-3	38607..38708	33	atcttcagtgtaaaatcgacagcca	atg	tag
101136	96ORF404	-3	21288..21389	33	cagacaccgtcttaagtcctttag	ata	taa

Table 11

SEQUENCE INFORMATION FOR PHAGES MATCHING WITH TABLE 1

M32695

Bacteriophage PM2 nuclease cleavage site

gi|166145|gb|M32695|BM2NCS [166145]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M32693

Bacteriophage PM2 Hind III fragment 4

gi|166144|gb|M32693|BM24HIND3 [166144]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M32693

Bacteriophage PM2 Hind III fragment 4

gi|166144|gb|M32693|BM24HIND3 [166144]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M32694

Bacteriophage PM2 Hind III fragment 3

gi|166143|gb|M32694|BM23HIND3 [166143]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

M26134

Bacteriophage PM2 structural protein gene containing purine/pyrimidine rich regions and anti-Z-DNA-IgG binding regions, complete cds

gi|289360|gb|M26134|BM2PROTIV [289360]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

J02452

bacteriophage fi 3'-terminal region rna

gi|215409|gb|J02452|PFITR3 [215409]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

AF020798

Bacteriophage Chp1 genome DNA, complete sequence

gi|217761|dbj|D00624|BCP1 [217761]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 12 protein links, or 1 genome link)

X72793

Clostridium botulinum C phage BONT/C1, ANTP-139, ANTP-33, ANTP-17, ANTP-70 genes and ORF-22

gi|516171|emb|X72793|CBCBONT [516171]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 6 protein links, or 4 nucleotide neighbors)

X51464

Clostridium botulinum D Phage C3 gene for exoenzyme C3

gi|14907|emb|X51464|CBDPE3 [14907]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

D90210

Bacteriophage c-st (from C. botulinum) C1-tox gene for botulinum C1 neurotoxin

gi|217780|dbj|D90210|CSTC1TOX [217780]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

S49407

type D neurotoxin [bacteriophage d-16 phi, host = C. botulinum, type D, CB16, Genomic, 4087 nt]
gi|260238|gb|S49407|S49407 [260238]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

X53370

Bacteriophage phi29 temperature sensitive mutant TS2(98) DNA polymerase gene
gi|15733|emb|X53370|POTS298 [15733]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)

X53371

Bacteriophage phi29 temperature sensitive mutant TS2(24) DNA polymerase gene
gi|15731|emb|X53371|POTS224 [15731]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 7 nucleotide neighbors)

X05973

Bacteriophage phi29 prohead RNA
gi|15680|emb|X05973|POP29PRO [15680]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 nucleotide neighbors)

V01155

Left end of bacteriophage phi-29 coding for 15 potential proteins Among
these are the terminal protein and the proteins encoded by the genes 1, 2 (sus), 3, and (probably) 4
gi|15659|emb|V01155|POP29B [15659]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 16 protein links, or 16 nucleotide neighbors)

X73097

Bacteriophage phi-29 left origin of replication
gi|312194|emb|X73097|BP29ORL [312194]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 5 nucleotide neighbors)

M14430

Bacteriophage phi-29 gene-17 gene, complete cds
gi|215321|gb|M14430|P29G17A [215321]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 8 nucleotide neighbors)

M14431

Bacteriophage phi-29 gene-16 gene, complete cds
gi|215319|gb|M14431|P29G16A [215319]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 7 nucleotide neighbors)

M20693

Bacteriophage phi-29 DNA, 3' end
gi|215343|gb|M20693|P29REPINB [215343]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 nucleotide neighbors)

M21016

Bacteriophage phi-29 DNA, 5' end
gi|215342|gb|M21016|P29REPINA [215342]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M12456

Bacteriophage phi-29 genes 9, 10 and 11 encoding p9 tail, incomplete, p10 connector, complete, and p11 lower collar, incomplete, respectively
gi|215338|gb|M12456|P29P9 [215338]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

M14782

Bacillus phage phi-29 head morphogenesis, major head protein, head fiber protein, tail protein, upper collar protein, lower collar protein, pre-neck appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds
gi|215323|gb|M14782|P29LATE2 [215323]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)

M26968

Bacteriophage phi-29 (from Bacillus subtilis) proteins p1 delta-1 genes, complete cds, and the sus1(629) mutation
gi|341558|gb|M26968|P29P1D1A [341558]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

J02448

Bacteriophage f1, complete genome
gi|166201|gb|J02448|F1CCG [166201]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, 205 nucleotide neighbors, or 1 genome link)

M24832

Bacteriophage f2 coat protein gene, partial cds
gi|166228|gb|M24832|F2CRNACA [166228]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

J02451

Bacteriophage fd, strain 478, complete genome
gi|215394|gb|J02451|PFDCG [215394]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 MEDLINE links, 10 protein links, 204 nucleotide neighbors, or 1 genome link)

M34834

Bacteriophage fr replicase gene, 5' end
gi|166139|gb|M34834|BFRREGRA [166139]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 9 nucleotide neighbors)

M38325

Bacteriophage fr replicase gene, 5' end
gi|166137|gb|M38325|BFRREGR [166137]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 9 nucleotide neighbors)

M35063

Bacteriophage fr coat protein replicase cistron (R region) RNA
gi|166134|gb|M35063|BFRRCRRA [166134]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 3 nucleotide neighbors)

S66567

alpha-atrial natriuretic factor/coat protein=fusion polypeptide [human, bacteriophage fr, expression vector pFAN15, PlasmidSyntheticRecombinant, 510 nt]
gi|435742|gb|S66567|S66567 [435742]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 15 nucleotide neighbors)

X15031

Bacteriophage fr RNA genome

gi|15071|emb|X15031|LEBFRX [15071]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or 1 genome link)

U51233

Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region light chain (IgM) mRNA, partial cds

gi|1277150|gb|U51233|MMU51233 [1277150]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 1669 nucleotide neighbors)

U51232

Mus musculus neutralizing anti-RNA-bacteriophage fr immunoglobulin variable region heavy chain (IgM) mRNA, partial cds

gi|1277148|gb|U51232|MMU51232 [1277148]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 1073 nucleotide neighbors)

U02303

Bacteriophage If1, complete genome

gi|3676280|gb|U02303|B2U02303 [3676280]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,10 protein links, or 1 genome link)

V00604

Phage M13 genome

gi|14959|emb|V00604|TNM13X [14959]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, or 205 nucleotide neighbors)

A32252

Synthetic bacteriophage M13 protein III probe

gi|1567340|emb|A32252|A32252 [1567340]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

A32251

Synthetic bacteriophage M13 protein III probe

gi|1567339|emb|A32251|A32251 [1567339]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M12465

Bacteriophage M13 mp10 mutations in lac operon

gi|215210|gb|M12465|M13LACMUT [215210]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 215 nucleotide neighbors)

M24177

Synthetic Bacteriophage M13 (clone M13.SV.B12) SV40 early promoter region DNA

gi|209416|gb|M24177|SYNSVB12 [209416]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M24176

Synthetic Bacteriophage M13 (clone M13.SV.B11) SV40 early promoter region DNA

gi|209415|gb|M24176|SYNSVB11 [209415]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M24175

Synthetic Bacteriophage M13 (clone M13.SV.8) SV40 early promoter region DNA

gi|208806|gb|M24175|SYNM13SV8 [208806]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 242 nucleotide neighbors)

M19979

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207813|gb|M19979|SYN33M13M [207813]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 617 nucleotide neighbors)

M19565

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207808|gb|M19565|SYN33M13H [207808]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 567 nucleotide neighbors)

M19564

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207807|gb|M19564|SYN33M13G [207807]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 12 nucleotide neighbors)

M19563

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207806|gb|M19563|SYN33M13F [207806]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 262 nucleotide neighbors)

M19561

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207804|gb|M19561|SYN33M13D [207804]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 27 nucleotide neighbors)

M19560

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207803|gb|M19560|SYN33M13C [207803]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M19559

Synthetic hybrids; recombinant DNA from bacteriophage M13 and plasmid pHV33

gi|207802|gb|M19559|SYN33M13B [207802]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 227 nucleotide neighbors)

M10568

Bacteriophage M13 replicative form II, replication origin, specific nick location

gi|215220|gb|M10568|M13ORIB [215220]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 650 nucleotide neighbors)

M10910

Bacteriophage M13 gene II regulatory region and M13sj1 mutant

gi|215209|gb|M10910|M13IIREG [215209]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 72 nucleotide neighbors)

M38295

Bacteriophage M13 HaeIII restriction fragment DNA

gi|215208|gb|M38295|M13HAEIII [215208]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 67 nucleotide neighbors)

E02067

DNA encoding a part of Bacteriophage M13 tg 127
gi|2170311|dbj|E02067|E02067 [2170311]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

J02467

Bacteriophage MS2, complete genome
gi|215232|gb|J02467|MS2CG [215232]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 8 MEDLINE links, 4 protein links, 20 nucleotide neighbors, or 1 genome link)

AJ004950

Bacteriophage P1 ban gene
gi|3688226|emb|AJ011592|BP1011592 [3688226]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

U88974

Bacteriophage P1 structural lytic transglycosylase (orf47), pep44b (orf44b),
pep44a (orf44a), and pep43 (orf43) genes, complete cds; and pep42 (orf42) gene, partial cds
gi|2661099|gb|AF035607|AF035607 [2661099]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 1 nucleotide neighbor)

AJ000741

Bacteriophage P1 darA operon
gi|2462938|emb|AJ000741|BPAJ7641 [2462938]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors)

X01828

Bacteriophage P1 recombinase gene cin
gi|15133|emb|X01828|MYP1CIN [15133]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

X98146

Bacteriophage P1 DNA sequence around the Op88 operator
gi|1359513|emb|X98146|BP1OP88OP [1359513]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

S61175

immI operon: icd=cell division repressor, ant1=antirepressor (promoters
P51a, P51b) [bacteriophage P1, Genomic, 728 nt]
gi|385908|gb|S61175|S61175 [385908]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 nucleotide neighbors)

X87824

Bacteriophage P1 gene 26
gi|861164|emb|X87824|XXBP1G26 [861164]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

X15638

Phage P1 DNA for lytic replicon containing promoter P53 and two open reading frames
gi|15735|emb|X15638|PP1LREP [15735]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 24 nucleotide neighbors)

X17512

Bacteriophage P1 DNA for immunity region immI

gi|15479|emb|X17512|P1IMMUNITY [15479]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 nucleotide neighbors)

X16005

Bacteriophage P1 c1 gene for P1c1 repressor protein

gi|15477|emb|X16005|P1C1 [15477]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

X03453

Bacteriophage P1 cre gene for recombinase protein

gi|15135|emb|X03453|MYP1CRE [15135]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

X06561

Bacteriophage P1 c1 gene 5'-region

gi|15128|emb|X06561|MYP1C1 [15128]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 6 nucleotide neighbors)

V01534

Bacteriophage P1 genome fragment (IS2 insertion spot). This regions contains

four unidentified reading frames and is known as insertion hot spot for IS2 insertion sequences

gi|15118|emb|V01534|MYOVP1 [15118]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors)

X56951

Bacteriophage P1 gene10

gi|406728|emb|X56951|BPP1GP10 [406728]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor)

K02380

Bacteriophage P1 replication region including repA, parA, and parB genes and

incA, incB, and incC incompatibility determinants

gi|215652|gb|K02380|P1REP [215652]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 MEDLINE links, 4 protein links, or 8 nucleotide neighbors)

X87674

Bacteriophage P1 lydA & lydB genes

gi|974763|emb|X87674|BACP1LYD [974763]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

X87673

Bacteriophage P1 gene 17

gi|974761|emb|X87673|BACP117 [974761]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M16618

Bacteriophage P1 c1 repressor binding sites

gi|215600|gb|M16618|PP1C1 [215600]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

SEG_PP1CIN

Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
 gi|215607|gb|SEG_PP1CIN [215607]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

K03173

Bacteriophage P1 C invertible element, right end, and cixR recombination site
 gi|215606|gb|K03173|PP1CIN2 [215606]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

215605

Bacteriophage P1 cin gene encoding recombinase, cixL recombination site, and 5' end of C invertible element
 gi|215605|lc|X01828 [215605]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M25470

Bacteriophage P1 tail fiber protein gene, complete cds
 gi|341349|gb|M25470|PP1TFPR [341349]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

M34382

Bacteriophage P1 sim region proteins, complete cds
 gi|215661|gb|M34382|PP1SIM [215661]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

M81956

Bacteriophage P1 R protein (R) gene, complete cds
 gi|215658|gb|M81956|PP1RP [215658]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

M37080

Bacteriophage P1 mini-P1 plasmid origin of replication
 gi|215657|gb|M37080|PP1REPOR [215657]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 46 nucleotide neighbors)

M27041

Bacteriophage P1 ref gene, complete cds
 gi|215650|gb|M27041|PP1REF [215650]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

L01408

Bacteriophage P1 partition protein (parB) gene, 3' end
 gi|215642|gb|L01408|PP1PARB [215642]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 41 nucleotide neighbors)

SEG_PP1PAR

Bacteriophage miniplasmid P1 parA gene, 5' end
 gi|215639|gb|SEG_PP1PAR [215639]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 48 nucleotide neighbors)

M36425

Bacteriophage miniplasmid P1 parB gene, 3' end
 gi|215638|gb|M36425|PP1PAR2 [215638]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

M36424

Bacteriophage miniplasmid P1 parA gene, 5' end
gi|215637|gb|M36424|PP1PAR1 [215637]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M11129

Bacteriophage P1 miniplasmid origin of replication region
gi|215632|gb|M11129|PP1ORIM [215632]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 43 nucleotide neighbors)

M25414

Bacteriophage P1 c1 repressor binding site, operator 88 (Op88)
gi|215631|gb|M25414|PP1OP88A [215631]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

M25413

Bacteriophage P1 c1 repressor binding site, operator 68 (Op68)
gi|215630|gb|M25413|PP1OP68A [215630]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M25412

Bacteriophage P1 c1 repressor binding site, operator 21 (Op21)
gi|215629|gb|M25412|PP1OP21A [215629]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M10510

Bacteriophage P1 recombination site loxR
gi|215628|gb|M10510|PP1LOXR [215628]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M10287

Bacteriophage P1 loxP X loxP recombination site
gi|215627|gb|M10287|PP1LOXPX [215627]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 nucleotide neighbors)

M10494

Bacteriophage P1 recombination site loxP
gi|215626|gb|M10494|PP1LOXP [215626]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 134 nucleotide neighbors)

M10511

Bacteriophage P1 recombination site loxL
gi|215625|gb|M10511|PP1LOXL [215625]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

M10512

Bacteriophage P1 recombination site loxB
gi|215624|gb|M10512|PP1LOXB [215624]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M10145

Bacteriophage P1 genome fragment with recombination site loxP
gi|215623|gb|M10145|PP1CREX [215623]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 21 nucleotide neighbors)

M13327

Bacteriophage P1 *Cin* recombinase activated cross over site, junction IV, clone pSHI326
gi|215622|gb|M13327|PP1CN26IV [215622]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13325

Bacteriophage P1 *Cin* recombinase activated cross over site, junction II, clone pSHI326
gi|215621|gb|M13325|PP1CN26II [215621]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1401 nucleotide neighbors)

M13323

Bacteriophage P1 *Cin* recombinase activated cross over site, junction IV, clone pSHI325
gi|215620|gb|M13323|PP1CN25IV [215620]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13321

Bacteriophage P1 *Cin* recombinase activated cross over site, junction II, clone pSHI325
gi|215619|gb|M13321|PP1CN25II [215619]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1058 nucleotide neighbors)

M13324

Bacteriophage P1 *Cin* recombinase activated cross over site, junction I, clone pSHI326
gi|215618|gb|M13324|PP1CIR26I [215618]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13319

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI327
gi|215617|gb|M13319|PP1CIN27R [215617]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13320

Bacteriophage P1 *Cin* recombinase activated cross over site, junction I, clone pSHI325
gi|215616|gb|M13320|PP1CIN25I [215616]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13318

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI324
gi|215615|gb|M13318|PP1CIN24L [215615]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1370 nucleotide neighbors)

M13317

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI323
gi|215614|gb|M13317|PP1CIN23M [215614]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1055 nucleotide neighbors)

M13316

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI323
gi|215613|gb|M13316|PP1CIN23L [215613]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13315

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI322
gi|215612|gb|M13315|PP1CIN22R [215612]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13314

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI322

gi|215611|gb|M13314|PP1CIN22L [215611]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1401 nucleotide neighbors)

M13313

Bacteriophage P1 *Cin* recombinase activated cross over site, right junction, clone pSHI321

gi|215610|gb|M13313|PP1CIN21R [215610]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 7 nucleotide neighbors)

M13312

Bacteriophage P1 *Cin* recombinase activated cross over site, left junction, clone pSHI321

gi|215609|gb|M13312|PP1CIN21L [215609]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1058 nucleotide neighbors)

M16568

Bacteriophage P1 *c4* repressor gene, complete cds

gi|215603|gb|M16568|PP1C4 [215603]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M13326

Bacteriophage P1 *Cin* recombinase activated cross over site, junction III, clone pSHI326

gi|215602|gb|M13326|PP1C26III [215602]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1192 nucleotide neighbors)

M13322

Bacteriophage P1 *Cin* recombinase activated cross over site, junction III, clone pSHI325

gi|215601|gb|M13322|PP1C25III [215601]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1231 nucleotide neighbors)

J05651

Bacteriophage P1 modulator protein (*bof*) gene, complete cds

gi|215598|gb|J05651|PP1BOFY1 [215598]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M33224

Bacteriophage P1 regulatory protein (*bof*) gene, complete cds

gi|215596|gb|M33224|PP1BOFFO [215596]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M10288

E.coli/bacteriophage P1 *loxR* recombination site

gi|146647|gb|M10288|ECOLOXR [146647]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

M10289

E.coli/bacteriophage P1 *loxL* recombination site

gi|146646|gb|M10289|ECOLOXL [146646]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M10290

E.coli *loxB* site, which can recombine with bacteriophage P1 *loxP* site

gi|146645|gb|M10290|ECOLOXB [146645]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M10287

Bacteriophage P1 loxP X loxP recombination site

gi|215627|gb|M10287|PP1LOXPX [215627]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 nucleotide neighbors)

M74046

Bacteriophage P1 pacA and pacB genes, complete cds

gi|215634|gb|M74046|PP1PACAB [215634]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 protein links)

M95666

Bacteriophage P1 gene 10, doc and phd genes, complete cds

gi|463276|gb|M95666|PP1PHDDOC [463276]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 4 protein links, or 1 nucleotide neighbor)

M25604

Bacteriophage Q-beta mutated autonomously replicating sequence MDV1 RNA fragment

gi|556359|gb|M25604|PQBARSMT [556359]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 8 nucleotide neighbors)

V00643

first half of the phage Q-beta gene for coat protein

gi|15088|emb|V00643|LEQBET [15088]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M25167

Bacteriophage Q-beta RNA fragment recovered from replicase binding complex

gi|556362|gb|M25167|PQBREPLICB [556362]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M24876

Bacteriophage Q-beta replicase RNA, 5' end

gi|556360|gb|M24876|PQBREPLICA [556360]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M25444

Synthetic bacteriophage Q-beta DNA

gi|209118|gb|M25444|SYNPQBTERM [209118]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 8 nucleotide neighbors)

M25463

Bacteriophage Q-beta self-replicating microvariant (+) RNA

gi|532489|gb|M25463|PQBMVSRRNA [532489]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M25014

Bacteriophage Q-beta RNA replicase gene, 5' end, and maturation protein gene, 3' end

gi|294316|gb|M25014|PQBREPLC [294316]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M25065

Bacteriophage Q-beta RNA sequence with putative stem loop

gi|294315|gb|M25065|PQBLOOP [294315]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

M10265

Bacteriophage Q-beta RNA molecule with the ability to replicate extracellularly

gi|215726|gb|M10265|PQBRNA [215726]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 8 nucleotide neighbors)

M24815

Bacteriophage Q-beta specified replicase subunit RNA,

gi|215725|gb|M24815|PQBREPL [215725]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 nucleotide neighbors)

M25461

Bacteriophage Q-beta plus-strand RNA, 5' terminus

gi|215724|gb|M25461|PQBPS5E [215724]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M25462

Bacteriophage Q-beta plus-strand RNA, 3' terminus

gi|215723|gb|M25462|PQBPS3E [215723]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 8 nucleotide neighbors)

M24871

Bacteriophage Q-beta nanovariant WSIII RNA

gi|215722|gb|M24871|PQBNVWSIC [215722]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M24870

Bacteriophage Q-beta nanovariant WSII RNA

gi|215721|gb|M24870|PQBNVWSIB [215721]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M24869

Bacteriophage Q-beta nanovariant WSI RNA

gi|215720|gb|M24869|PQBNVWSIA [215720]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M10495

Coliphage Q-beta MDV-1(+) RNA

gi|215719|gb|M10495|PQBMDV1A [215719]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 10 nucleotide neighbors)

J02484

bacteriophage qbeta coat protein cistron first half

gi|215717|gb|J02484|PQBCP5 [215717]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

M57754

Bacteriophage Q-beta minus strand RNA, 5' terminus

gi|215716|gb|M57754|PQBBMS5E [215716]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 8 nucleotide neighbors)

M24297

Bacteriophage Q-beta 5'-terminal region of the minus strand

gi|215715|gb|M24297|PQB5END [215715]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 8 nucleotide neighbors)

M10695

218

Bacteriophage Q-beta, MDV-1 RNA
gi|215714|gb|M10695|PQB1IR [215714]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 12 nucleotide neighbors)

M24827

Bacteriophage R17 A protein gene, 5' end
gi|216078|gb|M24827|R17RNACIS [216078]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)

M24829

Bacteriophage R17 coat protein gene, 5' end
gi|216075|gb|M24829|R17CP5 [216075]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)

J02488

bacteriophage r17 rna synthetase initiation site
gi|216080|gb|J02488|R17RNASYN [216080]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 2 protein links, or 6 nucleotide neighbors)

J02487

bacteriophage r17 coat protein initiation site
gi|216073|gb|J02487|R17COATP [216073]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

J02486

bacteriophage r17 a protein initiation site
gi|216071|gb|J02486|R17APROT [216071]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

M24826

Bacteriophage R17 coat protein RNA fragment
gi|216077|gb|M24826|R17CPRAA [216077]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 7 nucleotide neighbors)

M24296

Bacteriophage R17 3'-terminal fragment A RNA
gi|216070|gb|M24296|R173TFA [216070]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 9 nucleotide neighbors)

1TFN

structure refinement for a 24-nucleotide rna hairpin, nmr, minimized average
structure ribonucleic acid, hairpin, bacteriophage r17 mol_id: 1; molecule: r17c; chain: null; engineered: yes
gi|1942336|pdb|1TFN| [1942336]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 structure link)

1RPEA

rna (5'-d(gpgpgp apcpupg papcp g papupcp apcp g pcpapgpupcpupapu-3') (24-mer rna
hairpin coat protein binding site for bacteriophage r17) (nmr, minimized average structure)
gi|1421020|pdb|1RHT| [1421020]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 structure link)

M14428

Bacteriophage S13 circular DNA, complete genome

gi|216089|gb|M14428|S13CG [216089]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 12 protein links, 26 nucleotide neighbors, or 1 genome link)

J05393

Bacteriophage T1 DNA N-6-adenine-methyltransferase (M.T1) gene, complete cds

gi|166163|gb|J05393|BT1NAMTA [166163]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 protein links)

L46845

Bacteriophage T2 frd3, frd2 genes, complete cds

gi|951387|gb|L46845|PT2FRD32G [951387]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 17 nucleotide neighbors)

L43611

Bacteriophage T2 fibrin (wac) gene, complete cds

gi|903869|gb|L43611|PT2WAC [903869]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 4 nucleotide neighbors)

M24812

Bacteriophage T2 secondary structure RNA sequence

gi|215796|gb|M24812|PT2RNA [215796]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 nucleotide neighbors)

M22342

Bacteriophage T2 DNA-(adenine-N6)methyltransferase (dam) gene, complete cds

gi|215792|gb|M22342|PT2DAM [215792]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

S57515

orf 61.2 {intergenic region between 41 and 61} [bacteriophage T2, Genomic, 323 nt]

gi|298524|gb|S57515|S57515 [298524]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

X05312

Bacteriophage T2 gene 38 for receptor recognizing protein

gi|15197|emb|X05312|MYT2G38 [15197]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

X04442

Bacteriophage T2 gene 37 for receptor recognizing protein

gi|15195|emb|X04442|MYT2G37 [15195]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

X12460

Bacteriophage T2 gene 32 mRNA for single-stranded DNA binding protein

gi|15192|emb|X12460|MYT2G32 [15192]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 14 nucleotide neighbors)

X57797

Bacteriophage T2 gene for gp12

gi|14875|emb|X56555|BT2GP12 [14875]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 2 nucleotide neighbors)

X01755

Bacteriophage T2 tail fiber gene 36

gi|15189|emb|X01755|MYT2F36 [15189]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

M14784

Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis protein and DNA packaging proteins, complete cds

gi|215810|gb|M14784|PT3RE [215810]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, or 10 nucleotide neighbors)

SEG_PT3RNAPOL

Bacteriophage T3 RNA polymerase III gene, 5' end

gi|710559|gb|SEG_PT3RNAPOL [710559]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M22610

Bacteriophage T3 RNA polymerase III gene, 3' end

gi|340722|gb|M22610|PT3RNAPOL2 [340722]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M22609

Bacteriophage T3 RNA polymerase III gene, 5' end

gi|340721|gb|M22609|PT3RNAPOL1 [340721]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

X05031

Bacteriophage T3 gene region 1-2.5 with primary origin of replication

gi|15719|emb|X05031|POT3ORI [15719]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 5 nucleotide neighbors)

X03964

Bacteriophage T3 early control region pos. 308-810 from genome left end

gi|15718|emb|X03964|POT3EP [15718]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 20 nucleotide neighbors)

X17255

Bacteriophage T3 gene 1 to gene 11

gi|15682|emb|X17255|POT3111G [15682]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE links, 36 protein links, 17 nucleotide neighbors, or 1 genome link)

X15840

Phage T3 gene 10

gi|15625|emb|X15840|PODT3G10 [15625]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

X02981

Bacteriophage T3 gene 1 for RNA polymerase

gi|15561|emb|X02981|PODOT3P [15561]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

J02503

bacteriophage t3 5' end, terminally redundant sequence (trs)

gi|215816|gb|J02503|PT3TRS1 [215816]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

SEG_PT3TRS

bacteriophage t3 5' end, terminally redundant sequence (trs)

gi|215818|gb|SEG_PT3TRS [215818]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

J02504

bacteriophage t3 3' end, terminally redundant sequence (trs)

gi|215817|gb|J02504|PT3TRS2 [215817]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

HYPERLINK <http://www.rs.noda.sut.ac.jp/~kunisawa> h t t p://www.rs.noda.sut.ac.jp/~kunisawa
Bacteriophage T4 genomic database compiled by Arisaka et al.

X95646

Bacteriophage T5 DNA for region 60.5%-71% of the T5 genome

gi|2791557|emb|AJ001191|BTJ001191 [2791557]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,7 MEDLINE links, 12 protein links, or 6 nucleotide neighbors)

X56847

Bacteriophage T5 genomic region encoding early genes D10-D15

gi|15407|emb|X12930|MYT5D10 [15407]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 4 nucleotide neighbors)

AF039886

Bacteriophage T5 subclone T5.5.3r5.18r, single pass sequence, genomic survey sequence

gi|2811154|gb|AF039886|AF039886 [2811154]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039885

Bacteriophage T5 subclone T5.40f,41f, single pass sequence, genomic survey sequence

gi|2811153|gb|AF039885|AF039885 [2811153]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039884

Bacteriophage T5 subclone T5.26.fr, single pass sequence, genomic survey sequence

gi|2811152|gb|AF039884|AF039884 [2811152]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039883

Bacteriophage T5 subclone 10-T5.5.7F, single pass sequence, genomic survey sequence

gi|2811151|gb|AF039883|AF039883 [2811151]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039882

Bacteriophage T5 subclone 41-T5.5.4BF, single pass sequence, genomic survey sequence

gi|2811150|gb|AF039882|AF039882 [2811150]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039881

Bacteriophage T5 subclone 39-T5.5.4aF, single pass sequence, genomic survey sequence

gi|2811149|gb|AF039881|AF039881 [2811149]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 nucleotide neighbor)

222

AF039880

Bacteriophage T5 subclone 19-T5.7.2r, single pass sequence, genomic survey sequence
gi|2811148|gb|AF039880|AF039880 [2811148]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039879

Bacteriophage T5 subclone 18-T5.7.2F, single pass sequence, genomic survey sequence
gi|2811147|gb|AF039879|AF039879 [2811147]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039878

Bacteriophage T5 subclone 11-T5.5.7R, single pass sequence, genomic survey sequence
gi|2811146|gb|AF039878|AF039878 [2811146]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2
nucleotide neighbors)

AF039877

Bacteriophage T5 subclone T5.4FR, single pass sequence, genomic survey sequence
gi|2811145|gb|AF039877|AF039877 [2811145]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039876

Bacteriophage T5 subclone 22-T5.16R, single pass sequence, genomic survey sequence
gi|2811144|gb|AF039876|AF039876 [2811144]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039875

Bacteriophage T5 subclone 21-T5.16R, single pass sequence, genomic survey sequence
gi|2811143|gb|AF039875|AF039875 [2811143]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039874

Bacteriophage T5 subclone 21-T5.16F, single pass sequence, genomic survey sequence
gi|2811142|gb|AF039874|AF039874 [2811142]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039873

Bacteriophage T5 subclone 09-T5.6F, single pass sequence, genomic survey sequence
gi|2811141|gb|AF039873|AF039873 [2811141]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039872

Bacteriophage T5 subclone 09-T5.6R, single pass sequence, genomic survey sequence
gi|2811140|gb|AF039872|AF039872 [2811140]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 nucleotide neighbors)

AF039871

Bacteriophage T5 subclone 04-T5.26.R, single pass sequence, genomic survey sequence
gi|2811139|gb|AF039871|AF039871 [2811139]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF039870

Bacteriophage T5 subclone 13-T5.42F, single pass sequence, genomic survey sequence
gi|2811138|gb|AF039870|AF039870 [2811138]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

X69460

Bacteriophage T5 ltf gene for L-shaped tail fibers
gi|15415|emb|X69460|MYT5LTF [15415]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 1 protein link, or 4 nucleotide neighbors)

X03402

Bacteriophage T5 D15 gene for 5' exonuclease
gi|15413|emb|X03402|MYT5EXOG [15413]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

Z11972

Bacteriophage T5 tRNA-Tyr, tRNA-Glu, tRNA-Trp, tRNA-Phe, tRNA-Cys and
tRNA-Asn genes, and ORFs 91aa, 90aa, 42aa and 172aa
gi|15795|emb|Z11972|T56TRNAG [15795]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 3 nucleotide neighbors)

X03898

Bacteriophage T5 genes for tRNA-His, -Ser and -Leu
gi|15786|emb|X03898|STT5RN1 [15786]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 MEDLINE links)

X04177

Bacteriophage T5 gene for transfer RNA-Gln
gi|15421|emb|X04177|MYT5TRNQ [15421]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

X03899

Bacteriophage T5 genes for tRNA-Val, -Lys, -fMet, -Pro and -Ile3
gi|15787|emb|X03899|STT5RN2 [15787]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X03798

Bacteriophage T5 gene for tRNA-Asp (GUC)
gi|15472|emb|X03798|NCT5TRDG [15472]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

Y00364

Bacteriophage T5 tRNA gene cluster (27.8%-22.4%)
gi|15420|emb|Y00364|MYT5TRN [15420]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 nucleotide neighbors)

X03140

Bacteriophage T5 DNA with rho-dependent transcription terminator (Hind III-P fragment)
gi|15417|emb|X03140|MYT5RHO [15417]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

Z35070

Bacteriophage T6 DNA
gi|535228|emb|Z35074|MYEREGBT6 [535228]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

AF060870

Coliphage T6 small subunit distal tail fiber (gene 36) gene, partial cds; and large subunit distal tail fiber (gene 37) and tail fiber adhesin (gene 38) genes, complete cds

gi|3676458|gb|AF052605|AF052605 [3676458]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 protein links, or 2 nucleotide neighbors)

Z35072

Bacteriophage T6 DNA encoding ORF19.1 gene and g19 gene

gi|535232|emb|Z35072|MYTAILT6 [535232]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 protein links)

X12488

Bacteriophage T6 gene 32 mRNA for single-stranded DNA binding protein

gi|15843|emb|X12488|MYT6G32 [15843]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)

Z78095

Bacteriophage T6 DNA (1506 bp)

gi|1488562|emb|Z78095|BPHZ78095 [1488562]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 4 nucleotide neighbors)

Z35079

Bacteriophage T6 DNA for Ip5, Ip6

gi|535215|emb|Z35079|MY57BT6 [535215]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

X68725

E.coli bacteriophage T6 gene for beta-glucosyl-HMC-alpha-glucosyl-transferase

gi|296439|emb|X68725|ECT6 [296439]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

X69894

Bacteriophage T6 alt gene for ADP-Ribosyltransferase

gi|15422|emb|X69894|MYT6ADP [15422]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

L46846

Bacteriophage T6 frd3, frd2 genes, complete cds

gi|951390|gb|L46846|PT6FRD32G [951390]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

M27738

Bacteriophage T6 translational repressor protein (regA); complete cds

gi|215993|gb|M27738|PT6REGA [215993]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 5 nucleotide neighbors)

M38465

Bacteriophage T6 DNA ligase gene, complete cds

gi|215991|gb|M38465|PT6LIG55 [215991]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

V01146

Genome of bacteriophage T7

gi|431187|emb|V01146|T7CG [431187]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,13 MEDLINE links, 60 protein links, 105 nucleotide neighbors, or 1 genome link)

X60322

Bacteriophage alpha3 genes A, B, K, C, D, E, J, F, G, H

gi|14775|emb|X60322|BACALPHA [14775]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, 22 nucleotide neighbors, or 1 genome link)

X13332

Bacteriophage alpha3 DNA for origin of replication

gi|15093|emb|X13332|MIA3ORPL [15093]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X12611

Bacteriophage alpha3 gene for protein A part., finger domain

gi|15092|emb|X12611|MIA3AFIN [15092]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 6 nucleotide neighbors)

X15721

Bacteriophage alpha3 deletion mutation DNA for the origin region (-ori) of replication

gi|14774|emb|X15721|BA3DMOR9 [14774]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 11 nucleotide neighbors)

X15720

Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication

gi|14773|emb|X15720|BA3DMOR8 [14773]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)

X15719

Bacteriophage alpha3 insertion mutant DNA for the origin region (-ori) of replication

gi|14772|emb|X15719|BA3DMOR7 [14772]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 10 nucleotide neighbors)

X15718

Bacteriophage alpha3 deletion mutation DNA for origin region (-ori) of replication

gi|14771|emb|X15718|BA3DMOR6 [14771]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 11 nucleotide neighbors)

X15717

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication

gi|14770|emb|X15717|BA3DMOR5 [14770]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 9 nucleotide neighbors)

X15716

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication

gi|14769|emb|X15716|BA3DMOR4 [14769]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 10 nucleotide neighbors)

X15715

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
gi|14768|emb|X15715|BA3DMOR3 [14768]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)

X15714

Bacteriophage alpha3 deletion mutant DNA for origin region (-ori) of replication
gi|14767|emb|X15714|BA3DMOR2 [14767]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)

X15713

Bacteriophage alpha3 deletion mutant DNA for the origin region (-ori) of replication
gi|14766|emb|X15713|BA3DMOR1 [14766]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 11 nucleotide neighbors)

X62059

Bacteriophage alpha3 origin of cDNA synthesis (oriGA)
gi|14763|emb|X62059|AL3ORIGA [14763]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)

X62058

Bacteriophage alpha3 origin of cDNA synthesis (oriAA)
gi|14762|emb|X62058|AL3ORIAA [14762]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 13 nucleotide neighbors)

J02444

Bacteriophage alpha3 origin of DNA replication
gi|166103|gb|J02444|AL3ORI [166103]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

M25640

Bacteriophage alpha-3 H protein gene, complete cds
gi|166101|gb|M25640|AL3HP [166101]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)

M10631

Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
gi|166099|gb|M10631|AL3CSA [166099]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

X00774

Bacteriophage alpha-3 gene J sequence
gi|15431|emb|X00774|NCBA3J [15431]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

M25640

Bacteriophage alpha-3 H protein gene, complete cds
gi|166101|gb|M25640|AL3HP [166101]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 13 nucleotide neighbors)

M10631

Bacteriophage alpha-3 cleavage site for phage phi-X174 gene A protein
gi|166099|gb|M10631|AL3CSA [166099]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

J02459

Bacteriophage lambda, complete genome
gi|215104|gb|J02459|LAMCG [215104]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

J02482

Bacteriophage phi-X174, complete genome
gi|216019|gb|J02482|PX1CG [216019]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or 1 genome link)

J02454

Bacteriophage G4, complete genome
gi|215415|gb|J02454|PG4CG [215415]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 11 protein links, 20 nucleotide neighbors, or 1 genome link)

X60323

Bacteriophage phiK complete genome
gi|1478118|emb|X60323|BPHIKCG [1478118]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, 18 nucleotide neighbors, or 1 genome link)

L42820

Bacteriophage BF23 tail protein (hrs) gene, complete cds
gi|1048680|gb|L42820|BBFHRS [1048680]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X54455

Bacteriophage BF23 gene 17 and gene 18
gi|14797|emb|X54455|BF231718G [14797]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)

M37097

Bacteriophage BF23 DNA, right end of terminal repetition
gi|166115|gb|M37097|BBFRIGH [166115]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 nucleotide neighbors)

M37096

Bacteriophage BF23 DNA, left end of terminal repetition
gi|166114|gb|M37096|BBFLEFT [166114]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 nucleotide neighbor)

M37095

Bacteriophage BF23 A2-A3 gene, complete cds, and A1 gene, 5' end
gi|166110|gb|M37095|BBFA2A3 [166110]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 1 nucleotide neighbor)

AF056281

Bacteriophage BF23 clone bf23.mac5/6.1, genomic survey sequence
gi|3090930|gb|AF056281|AF056281 [3090930]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

AF056280

Bacteriophage BF23 clone bf23.mac3, genomic survey sequence
gi|3090929|gb|AF056280|AF056280 [3090929]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056279

Bacteriophage BF23 clone bf23.mac18/21.34, genomic survey sequence
gi|3090928|gb|AF056279|AF056279 [3090928]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056278

Bacteriophage BF23 clone bf23.mac16/19.33, genomic survey sequence
gi|3090927|gb|AF056278|AF056278 [3090927]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056277

Bacteriophage BF23 clone bf23.mac16/19-33, genomic survey sequence
gi|3090926|gb|AF056277|AF056277 [3090926]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056276

Bacteriophage BF23 clone bf23.mac12/9-9, genomic survey sequence
gi|3090925|gb|AF056276|AF056276 [3090925]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056275

Bacteriophage BF23 clone bf23.mac11/14-24, genomic survey sequence
gi|3090924|gb|AF056275|AF056275 [3090924]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056274

Bacteriophage BF23 clone bf23.57r64r, genomic survey sequence
gi|3090923|gb|AF056274|AF056274 [3090923]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 3 nucleotide neighbors)

AF056273

Bacteriophage BF23 clone bf23.54fr, genomic survey sequence
gi|3090922|gb|AF056273|AF056273 [3090922]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056272

Bacteriophage BF23 clone bf23.47fr.mac10/7, genomic survey sequence
gi|3090921|gb|AF056272|AF056272 [3090921]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056271

Bacteriophage BF23 clone bf23.23.66r, genomic survey sequence
gi|3090920|gb|AF056271|AF056271 [3090920]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056270

Bacteriophage BF23 clone bf23.23.64f, genomic survey sequence
gi|3090919|gb|AF056270|AF056270 [3090919]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056269

Bacteriophage BF23 clone bf23.23.60r, genomic survey sequence
gi|3090918|gb|AF056269|AF056269 [3090918]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056268

Bacteriophage BF23 clone bf23.23.60f, genomic survey sequence
gi|3090917|gb|AF056268|AF056268 [3090917]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 nucleotide neighbor)

AF056267

Bacteriophage BF23 clone bf23.23.59r, genomic survey sequence
gi|3090916|gb|AF056267|AF056267 [3090916]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056266

Bacteriophage BF23 clone bf23.23.59f, genomic survey sequence
gi|3090915|gb|AF056266|AF056266 [3090915]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056265

Bacteriophage BF23 clone bf23.23.56r, genomic survey sequence
gi|3090914|gb|AF056265|AF056265 [3090914]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056264

Bacteriophage BF23 clone bf23.23.56f, genomic survey sequence
gi|3090913|gb|AF056264|AF056264 [3090913]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056263

Bacteriophage BF23 clone bf23.23.68f55r, genomic survey sequence
gi|3090912|gb|AF056263|AF056263 [3090912]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056262

Bacteriophage BF23 clone bf23.23.43fr.66f, genomic survey sequence
gi|3090911|gb|AF056262|AF056262 [3090911]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056261

Bacteriophage BF23 clone bf23.23.2fr, genomic survey sequence
gi|3090910|gb|AF056261|AF056261 [3090910]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056260

Bacteriophage BF23 clone bf23.23.55.f, genomic survey sequence
gi|3090909|gb|AF056260|AF056260 [3090909]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056259

Bacteriophage BF23 clone bf23.23.53.r, genomic survey sequence
gi|3090908|gb|AF056259|AF056259 [3090908]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056258

Bacteriophage BF23 clone bf23.23.53.f, genomic survey sequence
gi|3090907|gb|AF056258|AF056258 [3090907]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056257

Bacteriophage BF23 clone bf23.23.52.r, genomic survey sequence
gi|3090906|gb|AF056257|AF056257 [3090906]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056256

Bacteriophage BF23 clone bf23.23.52.f, genomic survey sequence
gi|3090905|gb|AF056256|AF056256 [3090905]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056255

Bacteriophage BF23 clone bf23.23.49.r, genomic survey sequence
gi|3090904|gb|AF056255|AF056255 [3090904]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056254

Bacteriophage BF23 clone bf23.23.49.f, genomic survey sequence
gi|3090903|gb|AF056254|AF056254 [3090903]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056253

Bacteriophage BF23 clone bf23.23.48.r, genomic survey sequence
gi|3090902|gb|AF056253|AF056253 [3090902]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056252

Bacteriophage BF23 clone bf23.23.48.f, genomic survey sequence
gi|3090901|gb|AF056252|AF056252 [3090901]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056251

Bacteriophage BF23 clone bf23.23.44.r, genomic survey sequence
gi|3090900|gb|AF056251|AF056251 [3090900]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056250

Bacteriophage BF23 clone bf23.23.41.f, genomic survey sequence
gi|3090899|gb|AF056250|AF056250 [3090899]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056249

Bacteriophage BF23 clone bf23.23.22.a.r, genomic survey sequence
gi|3090898|gb|AF056249|AF056249 [3090898]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056248

Bacteriophage BF23 clone bf23.23.22.a.f, genomic survey sequence
gi|3090897|gb|AF056248|AF056248 [3090897]
(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

AF056247

Bacteriophage BF23 clone bf23.23.68.r, genomic survey sequence
gi|3090896|gb|AF056247|AF056247 [3090896]
(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

Z50114

Bacteriophage BF23 DNA for putative tail protein gene
gi|2464952|emb|Z50114|BF23LATE [2464952]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

D12824

Bacteriophage BF23 genes for minor tail protein gp24 and major tail protein gp25, complete cds
gi|520578|dbj|D12824|BBF2TAIL [520578]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

Z34953

Bacteriophage K3 ip9, ip7 and ip8 genes
gi|535261|emb|Z34953|MYK3IP978 [535261]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

Z35075

Bacteriophage K3 DNA for Ip3 and Ip4
gi|535229|emb|Z35075|MYEORF64K [535229]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

X05560

Bacteriophage K3 gene 38 for receptor recognizing protein
gi|15112|emb|X05560|MYK3G38 [15112]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

X04747

Bacteriophage K3 gene 37 for receptor recognizing protein
gi|15110|emb|X04747|MYK3G37 [15110]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

X01754

Bacteriophage K3 tail fiber gene 36
gi|15108|emb|X01754|MYK3F36 [15108]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

M16812

Bacteriophage K3 'r' lysis gene, complete cds
gi|215503|gb|M16812|PK3LYST [215503]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

L46833

Bacteriophage K3 frd3, frd2 genes, complete cds
gi|951377|gb|L46833|PK3FRD32G [951377]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 protein links, or 2 nucleotide neighbors)

L43613

Bacteriophage K3 fibrin (wac) gene, complete cds
gi|903861|gb|L43613|PK3WAC [903861]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 4 nucleotide neighbors)

X01753

Bacteriophage Ox2 tail fiber gene 36

gi|15122|emb|X01753|MYOX2F36 [15122]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

L43612

Bacteriophage Ox2 fibritin (wac) gene, complete cds

gi|903848|gb|L43612|OX2WAC [903848]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 4 nucleotide neighbors)

Z46880

Bacteriophage OX2 stp gene

gi|599663|emb|Z46880|BPOX2STP [599663]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

X05675

Bacteriophage Ox2 gene 38 for receptor-recognizing protein and flanking regions

gi|15124|emb|X05675|MYOX2G38 [15124]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

M33533

Bacteriophage RB18 translational repressor protein (regA) and Orf43.1, complete cds

gi|216083|gb|M33533|RB18REGA [216083]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

AF033329

Bacteriophage RB18 single-stranded binding protein (gene 32) gene, partial cds, and 5' region

gi|2645788|gb|AF033329|AF033329 [2645788]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 11 nucleotide neighbors)

M86231

Bacteriophage RB69 gene 62, 3'end; RegA (regA) gene, complete cds

gi|215354|gb|M86231|P6962REGA [215354]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

AF033332

Bacteriophage RB69 single-stranded binding protein (gene 32) gene, partial cds, and 5' region

gi|2645794|gb|AF033332|AF033332 [2645794]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 12 nucleotide neighbors)

U34036

Bacteriophage RB69 DNA polymerase (43) gene, complete cds

gi|1237125|gb|U34036|BRU34036 [1237125]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

V01145

Bacteriophage H1 genome fragment Each Thymine given in this sequence represents a HMU-residue

(HMU = 5-hydroxymethyluracil)

gi|15557|emb|V01145|PODOH1 [15557]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X05676

Bacteriophage M1 gene 38 for receptor recognizing protein and flanking regions

gi|15114|emb|X05676|MYM1G38 [15114]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

AF034575

Bacteriophage M1 putative integrase (int) gene, complete cds, and attP region, complete sequence
gi|2662472|gb|AF034575|AF034575 [2662472]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF033321

Bacteriophage M1 single-stranded binding protein (gene 32) gene, partial cds, and 5' region
gi|2645772|gb|AF033321|AF033321 [2645772]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)

X55190

Bacteriophage Tu1a 37 and 38 genes for receptor-recognizing proteins 37 and 38 (respectively), partial cds
gi|14860|emb|X55190|BPTU1A [14860]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

AF033334

Bacteriophage Tu1b single-stranded binding protein (gene 32) gene, partial cds, and 5' region
gi|2645798|gb|AF033334|AF033334 [2645798]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 5 nucleotide neighbors)

X55191

Bacteriophage Tu1b 37 gene for receptor-recognizing protein 37 (partial cds), 38 gene for receptor-recognizing protein 38, and t gene (partial cds)
gi|14863|emb|X55191|BPTU1B [14863]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

X13065

Bacteriophage phi80 early region
gi|14800|emb|X13065|BP80ER [14800]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 6 nucleotide neighbors)

D00360

Bacteriophage phi80 cor gene
gi|217782|dbj|D00360|P8080COR [217782]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 protein link)

X01639

Bacteriophage phi 80 DNA-fragment with replication origin
gi|15828|emb|X01639|XXPHI80 [15828]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 25 nucleotide neighbors)

X04051

Lambdoid bacteriophage phi 80 int-xis region (integrase-excisionase region)
gi|15770|emb|X04051|STPHI80X [15770]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

X06751

Phage Phi80 DNA for major coat protein
gi|15768|emb|X06751|STPHI80C [15768]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 11 nucleotide neighbors)

X75949

Bacteriophage phi80 DNA for ORF x171.8 and ORF x171.28'
gi|458811|emb|X75949|ECORF171B [458811]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 28 nucleotide neighbors)

- L40418
Bacteriophage phi-80 gene, complete cds
gi|1019107|gb|L40418|P80A [1019107]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)
- M24831
Bacteriophage phi-80 Tyr-tRNA gene, 3' end
gi|215363|gb|M24831|P80TGY [215363]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 43 nucleotide neighbors)
- M10670
Bacteriophage phi-80 replication origin
gi|215361|gb|M10670|P80ORI [215361]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)
- M24825
Bacteriophage phi-80 RNA fragment
gi|215360|gb|M24825|P80M3A [215360]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)
- M11919
Bacteriophage phi-80 cI immunity region encoding the N gene
gi|215358|gb|M11919|P80CI [215358]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)
- M10891
Bacteriophage phi-80 attP site DNA
gi|215357|gb|M10891|P80ATT [215357]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 nucleotide neighbor)
- M19473
Bacteriophage 933J (from E.coli) proviral Shiga-like toxin type I subunits A and B genes, complete cds
gi|215072|gb|M19473|J93SLTI [215072]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 2 protein links, or 20 nucleotide neighbors)
- Y10775
Bacteriophage 933W ileX, stx2A and stx2B genes
gi|1938206|emb|Y10775|BP933ILEX [1938206]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 36 nucleotide neighbors)
- X83722
Bacteriophage 933W slt-IIB gene
gi|1490229|emb|X83722|B933WSLT [1490229]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 20 nucleotide neighbors)
- X07865
Bacteriophage 933W slt-II gene for Shiga-like toxin typeII subunit A and B
gi|14892|emb|X07865|BWSLTII [14892]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 29 nucleotide neighbors)
- M16625
Bacteriophage H19B (from E.coli) sltIA and sltIB genes encoding Shiga-like toxin I subunits A and B, complete cds
gi|215043|gb|M16625|H19BSLT [215043]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 24 nucleotide neighbors)

M17358

Bacteriophage H19B shiga-like toxin-1 (SLT-1) A and B subunit DNA, complete cds

gi|215046|gb|M17358|H19BSLTA [215046]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 20 nucleotide neighbors)

U29728

Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds

gi|939708|gb|U29728|BNU29728 [939708]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 1 protein link)

J02580

Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (lc) and ORF1 genes. complete cds

gi|215366|gb|J02580|PA2LC [215366]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors)

U32222

Bacteriophage 186, complete sequence

gi|3337249|gb|U32222|B1U32222 [3337249]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

X51522

Bacteriophage P4 complete DNA genome

gi|450916|emb|X51522|MYP4CG [450916]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 13 protein links, 6 nucleotide neighbors. or 1 genome link)

X92588

Bacteriophage 82 orf33, orf151, orf56, orf96, rus, orf45, and Q genes

gi|1051111|emb|X92588|BAC82HOLL [1051111]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,7 protein links, or 1 nucleotide neighbor)

J02803

Bacteriophage 82 antitermination protein (Q) gene, complete cds

gi|215364|gb|J02803|P82Q [215364]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

U02466

Bacteriophage HK022 (cro), (cII) and (O) genes, complete cds, (P) gene, partial cds

gi|407285|gb|U02466|BHU02466 [407285]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

M26291

Bacteriophage D108 regulatory DNA-binding protein (ner) gene, complete cds

gi|166194|gb|M26291|D18NER [166194]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M11272

Bacteriophage D108 left-end DNA

gi|166193|gb|M11272|D18LEDNA [166193]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 2 nucleotide neighbors)

M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds

gi|166191|gb|M18902|D18KIL [166191]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

M10191

Bacteriophage D108, left end with Mu A protein binding sites L1 and L2

gi|166190|gb|M10191|D18BSL [166190]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)

J02447

bacteriophage d108 gene a 5' end

gi|166189|gb|J02447|D18AAA [166189]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 MEDLINE link)

V00865

Bacteriophage D108 fragment from genes A and ner (C-terminus of ner and N-terminus of A)

gi|15437|emb|V00865|NCD108 [15437]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

X01914

Bacteriophage IKe gene for DNA binding protein

gi|14957|emb|X01914|INIKEDBP [14957]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

AF064539

Bacteriophage N15, complete genome

gi|3192683|gb|AF064539|AF064539 [3192683]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

U02303

Bacteriophage If1, complete genome

gi|3676280|gb|U02303|B2U02303 [3676280]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 10 protein links, or 1 genome link)

AF007792

Bacteriophage Mu late morphogenetic region

gi|3551775|gb|AF007792|AF007792 [3551775]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

U24159

Bacteriophage HP1 strain HP1c1, complete genome

gi|1046235|gb|U24159|BHU24159 [1046235]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE links, 41 protein links, 8 nucleotide neighbors, or 1 genome link)

Z71579

Bacteriophage S2 type A 5.6 kb DNA fragment

gi|1679806|emb|Z71579|BPHS1ADNA [1679806]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 9 protein links, or 9 nucleotide neighbors)

X53238

Klebsiella sp. bacteriophage K11 gene 1 for RNA polymerase

gi|14984|emb|X53238|KSK11RPO [14984]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X85010

Bacteriophage A511 ply511 gene

gi|853748|emb|X85010|BPA511PLY [853748]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

U29728

Bacteriophage N4 single-stranded DNA-binding protein (N4SSB) gene, complete cds

gi|939708|gb|U29728|BNU29728 [939708]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 1 protein link)

J02445

bacteriophage bo1 3'-terminal region ma

gi|166152|gb|J02445|BO1TR3 [166152]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 5 nucleotide neighbors)

L06183Bacteriophage L5 (from *Leuconostoc oenos*) genome

gi|289353|gb|L06183|BL5GENM [289353]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 genome link)

AF074945

Mycoplasma arthritis bacteriophage MAV1, complete genome

gi|3511243|gb|AF074945|AF074945 [3511243]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 15 protein links, 3 nucleotide neighbors, or 1 genome link)

L13696Bacteriophage L2 (from *Mycoplasma*), complete genome

gi|289338|gb|L13696|BL2CG [289338]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 14 protein links, or 1 genome link)

X80191

Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins

gi|517237|emb|X80191|BPP7PR [517237]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 4 protein links, or 1 genome link)

M19377Bacteriophage Pf3 from *Pseudomonas aeruginosa* (New York strain), complete genome

gi|215380|gb|M19377|PF3COMNY [215380]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 5 nucleotide neighbors)

M11912Bacteriophage Pf3 from *Pseudomonas aeruginosa* (Nijmegen strain), complete genome

gi|215371|gb|M11912|PF3COMN [215371]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, 5 nucleotide neighbors, or 1 genome link)

V00605

Bacteriophage Pf1 gene encoding DNA binding protein

gi|14970|emb|V00605|INOPF1 [14970]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 1 nucleotide neighbor)

L05626

Bacteriophage PR4 capsid protein (P6) gene, complete cds

gi|215735|gb|L05626|PR4P6MAJA [215735]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

D13409

Bacteriophage phiCTX (isolated from *Pseudomonas aeruginosa*) cosR, attP, int genes
gi|217776|dbj|D13409|BPHCOSR [217776]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 3 nucleotide neighbors)

D13408

Bacteriophage phiCTX (isolated from *Pseudomonas aeruginosa*) cosL, ctx genes
gi|217775|dbj|D13408|BPHCOSLCTX [217775]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, or 3 nucleotide neighbors)

M24832

Bacteriophage f2 coat protein gene, partial cds
gi|166228|gb|M24832|F2CRNACA [166228]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618967|gb|AF017629|AF017629 [2618967]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds
gi|2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

AF017625

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618952|gb|AF017624|AF017624 [2618952]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618946|gb|AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017621

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618943|gb|AF017621|AF017621 [2618943]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

D26449

Bacteriophage PS17 FI gene for tail sheath protein (gpFI) and FII gene for tail tube protein (gpFII), complete cds

gi|452162|dbj|D26449|BPSFIFII [452162]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2 protein links)

X87627

Bacteriophage D3112 A and B genes

gi|974768|emb|X87627|BPD3112AB [974768]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

U32623

Bacteriophage D3 transcriptional activator CII (cII) gene, complete cds

gi|984852|gb|U32623|BDU32623 [984852]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 1 nucleotide neighbor)

L34781

Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and peptidoglycan hydrolase (lytA) gene, partial cds

gi|511838|gb|L34781|BPHHOLIN [511838]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors)

L14810

Bacteriophage P22 (gp10) gene, complete cds, and (gp26) gene, complete cds

gi|294053|gb|L14810|P22GP1026X [294053]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

X87420

Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators

gi|1143407|emb|X87420|BPES18GEN [1143407]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,5 protein links, or 9 nucleotide neighbors)

L42820

Bacteriophage BF23 tail protein (hrs) gene, complete cds

gi|1048680|gb|L42820|BBFHRS [1048680]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X14980

Bacteriophage PRD1 XV gene for protein P15 (lytic enzyme)

gi|15802|emb|X14980|TEPRD1XV [15802]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

X06321

Bacteriophage PRD1 gene 8 for DNA terminal protein

gi|15800|emb|X06321|TEPRD18 [15800]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 10 nucleotide neighbors)

X14336

Filamentous Bacteriophage I2-2 genome

gi|14920|emb|X14336|INBI22 [14920]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 9 protein links, 1 nucleotide neighbor, or 1 genome link)

L05001

240

Bacteriophage X glucosyl transferase gene, complete cds

gi|216044|gb|L05001|PXFCLUSYLT [216044]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

M29479

Bacteriophage p4 sid and psu genes partial cds, and delta gene, complete cds gi|215701|

gb|M29479|PP4SDP [215701]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 protein links, or 4 nucleotide neighbors)

SEG_PP4PSUSID

Bacteriophage P4 capsid size determination protein (sid) gene, 5' end

gi|215698|gb|SEG_PP4PSUSID [215698]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

M29650

Bacteriophage P4 polarity suppression protein (psu) gene, complete cds

gi|215697|gb|M29650|PP4PSUSID2 [215697]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M29651

Bacteriophage P4 capsid size determination protein (sid) gene, 5' end

gi|215696|gb|M29651|PP4PSUSID1 [215696]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M27748

Bacteriophage P4 gop, beta, and cII genes, complete cds and int gene, 3' end

gi|215691|gb|M27748|PP4GOPBC [215691]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 1 nucleotide neighbor)

K02750

Bacteriophage IKe, complete genome

gi|215061|gb|K02750|IKECG [215061]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 10 protein links, 4 nucleotide neighbors, or 1 genome link)

L40418

Bacteriophage phi-80 gene, complete cds

gi|1019107|gb|L40418|P80A [1019107]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

AF032122

Bacteriophage SfiI integrase (int) gene, partial cds; and bactoprenol glucosyl transferase (bgt), and glucosyl transferase II (grtII) genes, complete cds

gi|2465412|gb|AF021347|AF021347 [2465412]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 2 nucleotide neighbors)

M35825

Bacteriophage SF6 fragment D lysozyme gene, complete cds

gi|216105|gb|M35825|SF6LYZ [216105]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 protein link)

Z35479

Bacteriophage C16 ip1 gene

gi|534936|emb|Z35479|BC16IP1 [534936]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 2 nucleotide neighbors)

X12638

Bacteriophage 21 DNA for gene 2

gi|296141|emb|X12638|B21GENE2 [296141]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

X02501

Bacteriophage 21 DNA for left end sequence with genes 1 and 2

gi|15825|emb|X02501|XXPHA21 [15825]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds

gi|215466|gb|M65239|PH2LYSGEN [215466]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

M58702

Bacteriophage 21 late gene regulatory region

gi|215465|gb|M58702|PH2LATEGE [215465]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M81255

Bacteriophage 21 head gene operon

gi|215454|gb|M81255|PH2HEADTL [215454]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 10 protein links, or 4 nucleotide neighbors)

M23775

Bacteriophage 21 glycoprotein 1 gene, complete cds, and glycoprotein gene, 5' end

gi|215451|gb|M23775|PH2GPA [215451]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

M61865

Bacteriophage 21 excisionase (xis), integrase (int) and isocitrate dehydrogenase (icd), complete cds

gi|215448|gb|M61865|PH22XISAA [215448]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 9 nucleotide neighbors)

S72011

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618967|gb|AF017629|AF017629 [2618967]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017628

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618964|gb|AF017628|AF017628 [2618964]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017627

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds

gi|2618961|gb|AF017627|AF017627 [2618961]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017626

Bacteriophage 21 isocitrate dehydrogenase (icd) gene, partial cds; and integrase (int) gene, partial cds

gi|2618958|gb|AF017626|AF017626 [2618958]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 49 nucleotide neighbors)

AF017625

242
Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618955|gb|AF017625|AF017625 [2618955]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017624

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618952|gb|AF017624|AF017624 [2618952]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017623

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618949|gb|AF017623|AF017623 [2618949]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017622

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618946|gb|AF017622|AF017622 [2618946]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

AF017621

Bacteriophage 21 isocitrate dehydrogenase (icd) and integrase (int) genes, partial cds
gi|2618943|gb|AF017621|AF017621 [2618943]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 44 nucleotide neighbors)

M57455

Bacteriophage 42D (clone pDB17) (from *Staphylococcus aureus*) staphylokinase gene, complete cds
gi|215344|gb|M57455|P42STK [215344]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 9 nucleotide neighbors)

Y12633

Bacteriophage 85 DNA, promoter sequence of unknown gene
gi|2058285|emb|Y12633|B85PROM [2058285]

(View GenBank report, FASTA report, ASN.1 report, or Graphical view)

X98146

Bacteriophage P1 DNA sequence around the Op88 operator
gi|1359513|emb|X98146|BP1OP88OP [1359513]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 1 nucleotide neighbor)

Y07739

Staphylococcus phage Twort holTW, plyTW genes
gi|2764979|emb|Y07739|BPTWGHOLG [2764979]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2 protein links)

L07580

Bacteriophage phi-11 rinA and rinB genes, required for the activation of *Staphylococcal* phage phi-11 int expression
gi|166160|gb|L07580|BPHRINAB [166160]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 2 protein links)

M34832

Bacteriophage phi-11 integrase (int) and excisionase (xis) genes, complete cds
gi|166157|gb|M34832|BPHINTXIS [166157]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

M20394

Bacteriophage phi-11 S.aureus attachment site (attP)

gi|166156|gb|M20394|BPHATTTP [166156]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 4 nucleotide neighbors)

X23128

Bacteriophage phi-13 integrase gene

gi|758228|emb|X82312|PHI13INT [758228]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 protein link, or 3 nucleotide neighbors)

X61719

S.aureus phi-13 lysogen right chromosome/bacteriophage DNA junction

gi|46625|emb|X61719|SAP13RJNC [46625]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X61718

S.aureus phi-13 lysogen left chromosomal/bacteriophage DNA junction

gi|46624|emb|X61718|SAP13LJNC [46624]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

X61717

Bacteriophage phi-13 core sequence for attachment

gi|14799|emb|X61717|BP13ATTTP [14799]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 3 nucleotide neighbors)

U01875

Bacteriophage phi-13 putative regulatory region and integrase (int) gene, partial cds

gi|437118|gb|U01875|U01875 [437118]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, or 4 nucleotide neighbors)

X67739

S.aureus Bacteriophage phi-42 attP gene

gi|14809|emb|X67739|BPATTTPA [14809]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 nucleotide neighbors)

U01872

Bacteriophage phi-42 integrase (int) gene, complete cds

gi|437115|gb|U01872|U01872 [437115]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 2 protein links, or 3 nucleotide neighbors)

X94423

Staphylococcus aureus bacteriophage phi-42 DNA with ORFs (restriction modification system)

gi|1771597|emb|X94423|SARMS [1771597]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 1 nucleotide neighbor)

M27965

Bacteriophage L54a (from S.aureus) int and xis genes, complete cds

gi|215096|gb|M27965|L54INTXIS [215096]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, MEDLINE 1 link, 2 protein links, or 3 nucleotide neighbors)

U72397

Bacteriophage 80 alpha holin and amidase genes, complete cds

gi|1763241|gb|U72397|B8U72397 [1763241]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 2 nucleotide neighbors)

AB009866

Bacteriophage phi PVL proviral DNA, complete sequence

gi|3341907|dbj|AB009866|AB009866 [3341907]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, or 1 nucleotide neighbor)

Z47794

Bacteriophage Cp-1 DNA, complete genome

gi|2288892|emb|Z47794|BPCP1XX [2288892]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or 1 genome link)

SEG_CP7RSIT

Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat

gi|166186|gb|SEG_CP7RSIT [166186]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M11635

Bacteriophage Cp-7 (S.pneumoniae) DNA, 3' inverted terminal repeat

gi|166185|gb|M11635|CP7RSIT2 [166185]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M11636

Bacteriophage Cp-7 (S.pneumoniae) 5' inverted terminal repeat

gi|166184|gb|M11636|CP7RSIT1 [166184]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

SEG_CP5RSIT

Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat

gi|166181|gb|SEG_CP5RSIT [166181]

(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1 MEDLINE link)

M11633

Bacteriophage Cp-5 (S.pneumoniae) 3' inverted terminal repeat

gi|166180|gb|M11633|CP5RSIT2 [166180]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M11634

Bacteriophage Cp-5 (S.pneumoniae), 5' inverted terminal repeat

gi|166179|gb|M11634|CP5RSIT1 [166179]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

M34780

Bacteriophage Cp-9 muramidase (cp19) gene

gi|166187|gb|M34780|CP9CPL [166187]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 1 nucleotide neighbor)

M34652

Bacteriophage HB-3 amidase (hbl) gene, complete cds

gi|215055|gb|M34652|HB3HBLA [215055]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

U64984

Streptococcus pyogenes phage T12 repressor, excisionase (xis), integrase(int) and erythrogenic toxin A precursor (speA) genes, complete cds gi|1877426|gb|U40453|SPU40453 [1877426]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 4 protein links, or 22 nucleotide neighbors)

X12375

Phage CP-T1 (*Vibrio cholerae*) DNA for packaging signal (pac site)

gi|15435|emb|X12375|NCCPPAC [15435]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF087814

Vibrio cholerae filamentous bacteriophage fs-2 DNA, complete genome sequence

gi|3702207|dbj|AB002632|AB002632 [3702207]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 9 protein links, or 1 genome link)

D83518

Bacteriophage KVP40 gene for major capsid protein precursor, complete cds

gi|3046858|dbj|D83518|D83518 [3046858]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 1 protein link)

AF033322

Bacteriophage PST single-stranded binding protein (gene 32) gene, partial cds, and 5' region

gi|2645774|gb|AF033322|AF033322 [2645774]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 protein link, or 17 nucleotide neighbors)

X94331

Bacteriophage L cro, 24, c2, and c1 genes

gi|1469213|emb|X94331|BLCRO24C [1469213]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 protein links)

U82619

Shigella flexneri bacteriophage V glucosyl transferase (gtr), integrase (int) and excisionase (xis) genes, complete cds

gi|2465470|gb|U82619|SFU82619 [2465470]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 8 protein links, or 1 nucleotide neighbor)

246
Table 12

NCBI *Entrez* Nucleotide QUERY

Key words: bacteriophage and lysis

56 citations found (all selected)

AJ011581

Bacteriophage PS119 lysis genes 13, 19, 15, and packaging gene 3, complete cds
gi13676084|embl|AJ011581|BPS011581 [3676084]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

AJ011580

Bacteriophage PS34 lysis genes 13, 19, 15, antiterminator gene 23, and packaging gene 3, complete cds
gi13676078|embl|AJ011580|BPS011580 [3676078]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 2 nucleotide neighbors)

AJ011579

Bacteriophage PS3 lysis genes 13, 19, 15, and packaging gene 3
gi13676073|embl|AJ011579|BPS011579 [3676073]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 protein links, or 1 nucleotide neighbor)

AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cl protein (cl), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds
gi12668751|gb|AF034975 [2668751]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

U37314

Bacteriophage lambda Rz1 protein precursor (Rz1) gene, complete cds
gi1017780|gb|U37314|BLU37314 [1017780]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 1 protein link, or 9 nucleotide neighbors)

U00005

E. coli hflA locus encoding the hflX, hflK and hflC genes, hflq gene, complete cds; miaA gene, partial cds
gi1436153|gb|U00005|ECOHLA [436153]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 1 protein link, or 9 nucleotide neighbors)

links, 5 protein links, or 8 nucleotide neighbors)

U32222

Bacteriophage 186, complete sequence
 gil3337249|gb|U32222|B|U32222 [3337249]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 6 MEDLINE
 links, 46 protein links, or 5 nucleotide neighbors)

AF064539

Bacteriophage N15, complete genome
 gil3192683|gb|AF064539|AF064539 [3192683]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE
 links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

AF063097

Bacteriophage P2, complete genome
 gil3139086|gb|AF063097|AF063097 [3139086]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 21 MEDLINE
 links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes
 gil2707950|emb|Z97974|BPHIADH [2707950]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE
 links, 9 protein links, or 1 nucleotide neighbor)

AF059243

Bacteriophage NL95, complete genome
 gil3088545|gb|AF059243|AF059243 [3088545]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE
 links, 4 protein links, 3 nucleotide neighbors, or 1 genome link)

AF052431

Bacteriophage M11 A-protein, coat protein, A1-protein, and replicase
 genes, complete cds
 gil2981208|gb|AF052431 [2981208]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE
 links, 4 protein links, or 8 nucleotide neighbors)

Y07739

Staphylococcus phage Twort holTW, plyTW genes
 gil2764979|emb|Y07739|BPTWGHOLG [2764979]
 (View GenBank report, FASTA report, ASN.1 report, Graphical view, or 2
 protein links)

X94331

Bacteriophage L. cro, 24, c2, and c1 genes
gil1469213|emblX94331|BLCRO24C [1469213]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 4 protein links)

X78410

Bacteriophage phiadh holin and lysin genes
gil793848|emblX78410|LGHOLLYS [793848]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

X99260

Bacteriophage B103 genomic sequence
gil1429229|emblX99260|BB103G [1429229]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 17 protein links, or 12 nucleotide neighbors)

AJ000741

Bacteriophage P1 darA operon
gil2462938|emblAJ000741|BPAJ7641 [2462938]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 10 protein links, or 31 nucleotide neighbors)

X87420

Bacteriophage ES18 genes 24, c2, cro, c1, 18, and oL and oR operators
gil1143407|emblX87420|BPES18GEN [1143407]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 5 protein links, or 9 nucleotide neighbors)

L35561

Bacteriophage phi-105 ORFs 1-3
gil532218|gbL35561|PHSORFHTR [532218]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, or 3 protein links)

D10027

Group II RNA coliphage GA genome
gil217784|dbjD10027|PGAXX [217784]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, 5 nucleotide neighbors, or 1 genome link)

V01128

Bacteriophage phi-X174 (cs70 mutation) complete genome
gil15535|emblV01128|PHIX174 [15535]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 4 MEDLINE links, 11 protein links, or 26 nucleotide neighbors)

S81763

coat gene...replicase gene [bacteriophage KU1, host=Escherichia coli,
group II RNA phage, Genomic RNA, 3 genes, 120 nt]
gil1438766|gb|S81763|S81763 [1438766]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 1
MEDLINE link)

U38906

Bacteriophage r1t integrase, repressor protein (rro), dUTPase, holin and
lysin genes, complete cds
gil1353517|gb|U38906|BRU38906 [1353517]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, 50 protein links, or 3 nucleotide neighbors)

X91149

Bacteriophage phi-C31 DNA cos region
gil1107473|emb|X91149|APHIC31C [1107473]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 6 protein links, or 1 nucleotide neighbor)

V00642

phage MS2 genome
gil15081|emb|V00642|LEMS2X [15081]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE
links, 4 protein links, or 20 nucleotide neighbors)

V01146

Genome of bacteriophage T7
gil431187|emb|V01146|T7CG [431187]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,13 MEDLINE
links, 60 protein links, 105 nucleotide neighbors, or 1 genome link)

X78401

Bacteriophage P22 right operon, orf 48, replication genes 18 and 12, nin
region genes, ninG phosphatase, late control gene 23, orf 60, complete-
cds, late control region, start of lysis gene 13
gil512343|emb|X78401|POP22NIN [512343]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE
links, 13 protein links, or 4 nucleotide neighbors)

Y00408

Bacteriophage T4 gene t for lysis protein
gil15368|emb|Y00408|MYT4T [15368]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 1 protein link, or 3 nucleotide neighbors)

Z26590

Bacteriophage mv4 lysA and lysB genes
gil410500|emb|Z26590|MV4LYSAB [410500]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

X07809

Phage phiX174 lysis (E) gene upstream region
gil15094|emb|X07809|MIPHXE [15094]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 4 nucleotide neighbors)

Z34528

Lactococcal bacteriophage c2 lysin gene
gil506455|emb|Z34528|LBC2LYSIN [506455]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 4 nucleotide neighbors)

X15031

Bacteriophage fr RNA genome
gil15071|emb|X15031|LEBFRX [15071]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, 9 nucleotide neighbors, or 1 genome link)

X80191

Bacteriophage PP7 mRNA for maturation, coat, lysis and replicase proteins
gil517237|emb|X80191|BPP7PR [517237]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 1 genome link)

X85010

Bacteriophage A511 ply511 gene
gil853748|emb|X85010|BPA511PLY [853748]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

X85009

Bacteriophage A500 hol500 and ply500 genes
gil853744|emb|X85009|BPA500PLY [853744]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 4 nucleotide neighbors)

X85008

Bacteriophage A118 hol118 and ply118 genes
gil853740|emb|X85008|BPA118PLY [853740]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

251

Z35638

Bacteriophage phi-X174 genes for lysis protein and beta-lactamase
gil520996|embl|Z35638|BPLYSPR [520996]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE
link, 2 protein links, or 516 nucleotide neighbors)

J02459

Bacteriophage lambda, complete genome
gil215104|gb|J02459|LAMCG [215104]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 87 MEDLINE
links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

X87674

Bacteriophage P1 lydA & lydB genes
gil974763|embl|X87674|BACP1LYD [974763]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE
link, 2 protein links, or 2 nucleotide neighbors)

X87673

Bacteriophage P1 gene 17
gil974761|embl|X87673|BACP117 [974761]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE
link, 1 protein link, or 1 nucleotide neighbor)

M14784

Bacteriophage T3 strain amNG220B right end, tail fiber protein, lysis
protein and DNA packaging proteins, complete cds
gil215810|gb|M14784|PT3RE [215810]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE
link, 9 protein links, or 10 nucleotide neighbors)

M11813

Bacteriophage PZA (from B.subtilis), complete genome
gil216046|gb|M11813|PZACG [216046]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE
links, 27 protein links, 17 nucleotide neighbors, or 1 genome link)

M16812

Bacteriophage K3 't' lysis gene, complete cds
gil215503|gb|M16812|PK3LYST [215503]
(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE
link, 1 protein link, or 4 nucleotide neighbors)

J04356

Bacteriophage P22 proteins 15 (complete cds), and 19 (3' end) genes
gil215265|gb|J04356|P221SP [215265]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

J04343

Bacteriophage JP34 coat and lysis protein genes, complete cds, and replicase protein gene, 5' end
gil215076|gb|J04343|JP3COLY [215076]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 2 nucleotide neighbors)

J02482

Bacteriophage phi-X174, complete genome
gil216019|gb|J02482|PX1CG [216019]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,23 MEDLINE links, 11 protein links, 26 nucleotide neighbors, or 1 genome link)

M99441

Bacteriophage T4 anti-sigma 70 protein (asiA) gene, complete cds and lysis protein, 3' end
gil215820|gb|M99441|PT4ASIA [215820]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 2 protein links, or 2 nucleotide neighbors)

M65239

Bacteriophage 21 lysis genes S, R, and Rz, complete cds
gil215466|gb|M65239|PH2LYSGEN [215466]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

M10637

Phage G4 D/E overlapping gene system, encoding D (morphogenetic) and E (lysis) proteins
gil215427|gb|M10637|PG4DE [215427]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 12 nucleotide neighbors)

J02454

Bacteriophage G4, complete genome
gil215415|gb|J02454|PG4CG [215415]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 11 protein links, 20 nucleotide neighbors, or 1 genome link)

J02580

Bacteriophage PA-2 (E.coli porcine strain isolate) Rz gene, 5'end; ORF2, outer membrane porin protein (Ic) and ORF1 genes, complete cds
gil215366|gb|J02580|PA2LC [215366]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 4 protein links, or 4 nucleotide neighbors)

M14782

Bacillus phage phi-29 head morphogenesis, major head protein, head fiber protein, tail protein, upper collar protein, lower collar protein, pre-neck appendage protein, morphogenesis(13), lysis, morphogenesis(15), encapsidation genes, complete cds
gil215323|gb|M14782|P29LATE2 [215323]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 11 protein links, or 11 nucleotide neighbors)

M10997

Bacteriophage P22 lysis genes 13 and 19, complete cds
gil215262|gb|M10997|P221319 [215262]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 3 nucleotide neighbors)

J02467

Bacteriophage MS2, complete genome
gil215232|gb|J02467|MS2CG [215232]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,8 MEDLINE links, 4 protein links, 20 nucleotide neighbors, or 1 genome link)

M14035

Bacteriophage lambda lysis S gene with mutations leading to nonlethality of S in the plasmid pRG1
gil215180|gb|M14035|LAMLYS [215180]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 1 protein link, or 14 nucleotide neighbors)

U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein hydrolase (lysB) gene, complete cds
gil530796|gb|U04309|BPU04309 [530796]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Table 13

NCBI *Entrez* Nucleotide QUERY

Key word: holin

51 citations found (all selected)

AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds
 gil2668751|gb|AF034975| [2668751]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

U52961

Staphylococcus aureus holin-like protein LrgA (lrgA) and LrgB (lrgB) genes, complete cds
 gil1841516|gb|U52961|SAU52961 [1841516]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

U28154

Haemophilus somnus cryptic prophage genes, capsid scaffolding protein gene, partial cds, major capsid protein precursor, endonuclease, capsid completion protein, tail synthesis proteins, holin, and lysozyme genes, complete cds
 gil1765928|gb|U28154|HSU28154 [1765928]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 13 protein links)

AF032122

Streptococcus thermophilus bacteriophage Sfi19 central region of genome
 gil2935682|gb|AF032122| [2935682]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

AF032121

Streptococcus thermophilus bacteriophage Sfi21 central region of genome
 gil2935667|gb|AF032121|AF032121 [2935667]
 (View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 14 protein links, or 2 nucleotide neighbors)

AF021803

Bacillus subtilis 168 prophage SPbeta N-acetylmuramoyl-L-alanine amidase (blyA), holin-like protein (bhlA), holin-like protein (bhlB), and yolK genes, complete cds; and yolJ gene, partial cds
gi|2997594|gb|AF021803|AF021803 [2997594]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 5 protein links, or 1 nucleotide neighbor)

AF057033

Streptococcus thermophilus bacteriophage sfil1 gp502 (orf502), gp284 (orf284), gp129 (orf129), gp193 (orf193), gp119 (orf119), gp348 (orf348), gp53 (orf53), gp113 (orf113), gp104 (orf104), gp114 (orf114), gp128 (orf128), gp168 (orf168), gp117 (orf117), gp105 (orf105), putative minor tail protein (orf1510), putative minor structural protein (orf512), putative minor structural protein (orf1000), gp373 (orf373), gp57 (orf57), putative anti-receptor (orf695), putative minor structural protein (orf669), gp149 (orf149), putative holin (orf141), putative holin (orf87), and lysin (orf288) genes, complete cds
gi|3320432|gb|AF057033|AF057033 [3320432]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,25 protein links, or 1 nucleotide neighbor)

U32222

Bacteriophage 186, complete sequence
gi|3337249|gb|U32222|B1U32222 [3337249]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 46 protein links, or 5 nucleotide neighbors)

AB009866

Bacteriophage phi PVL proviral DNA, complete sequence
gi|3341907|dbj|AB009866|AB009866 [3341907]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, or 1 nucleotide neighbor)

AF009630

Bacteriophage bIL170, complete genome
gi|3282260|gb|AF009630|AF009630 [3282260]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,63 protein links, 3 nucleotide neighbors, or 1 genome link)

AF064539

Bacteriophage N15, complete genome

gil3192683|gb|AF064539|AF064539 [3192683]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 60 protein links, 26 nucleotide neighbors, or 1 genome link)

AF063097

Bacteriophage P2, complete genome

gil3139086|gb|AF063097|AF063097 [3139086]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,21 MEDLINE links, 42 protein links, 3 nucleotide neighbors, or 1 genome link)

Z97974

Bacteriophage phiadh lys, hol, intG, rad, and tec genes

gil2707950|emb|Z97974|BPHIADH [2707950]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 9 protein links, or 1 nucleotide neighbor)

X95646

Streptococcus thermophilus bacteriophage Sfi21 DNA; lysogeny module, 8141 bp

gil2292747|emb|X95646|BSFI21LYS [2292747]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 19 protein links, or 3 nucleotide neighbors)

SEG_LLHLYSIN0

Bacteriophage LL-H structural protein gene, partial cds; minor structural protein gp61 (g57), unknown protein, unknown protein, structural protein (g20), unknown protein, unknown protein, major capsid protein (g34), main tail protein gp19 (g17), holin (hol), muramidase (mur), unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, and unknown protein genes, complete cds; unknown protein gene, partial cds; and unknown protein, unknown protein, unknown protein, unknown protein, unknown protein, minor structural protein gp75 (g70), minor structural protein gp89 (g88), minor structural protein gp58 (g71), unknown protein, unknown protein, unknown protein, and unknown protein genes, complete cds

gil1004337|gb|SEG_LLHLYSIN0 [1004337]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 MEDLINE links, 31 protein links, or 1 nucleotide neighbor)

M96254

Bacteriophage LL-H holin (hol), muramidase (mur), and unknown protein genes, complete cds

gil1004336|gb|M96254|LLHLYSIN03 [1004336]

(View GenBank report,FASTA report,ASN.1 report, or Graphical view)

Y07740

Staphylococcus phage 187 ply187 and hol187 genes
gil2764982|embl|Y07740|BP187PLYH [2764982]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 2
protein links)

U88974

Streptococcus thermophilus bacteriophage 01205 DNA sequence
gil2444080|gb|U88974| [2444080]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 57 protein links, or 6 nucleotide neighbors)

Z99117

Bacillus subtilis complete genome (section 14 of 21): from 2599451 to
2812870
gil2634966|embl|Z99117|BSUB0014 [2634966]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,233
protein links, 51 nucleotide neighbors, or 1 genome link)

Z99115

Bacillus subtilis complete genome (section 12 of 21): from 2195541 to
2409220
gil2634478|embl|Z99115|BSUB0012 [2634478]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,244
protein links, 64 nucleotide neighbors, or 1 genome link)

Z99110

Bacillus subtilis complete genome (section 7 of 21): from 1194391 to
1411140
gil2633472|embl|Z99110|BSUB0007 [2633472]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,226
protein links, 31 nucleotide neighbors, or 1 genome link)

X78410

Bacteriophage phiadh holin and lysin genes
gil793848|embl|X78410|LGHOLLYS [793848]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 2 protein links, or 1 nucleotide neighbor)

Z93946

Bacteriophage Dp-1 dph and pal genes and 5 open reading frames
gil1934760|emblZ93946|BPDP1ORFS [1934760]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 6 protein links)

AF011378

Bacteriophage sk1 complete genome
gil2392824|gb|AF011378|AF011378 [2392824]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,54 protein links, 2 nucleotide neighbors, or 1 genome link)

Z47794

Bacteriophage Cp-1 DNA, complete genome
gil2288892|emblZ47794|BPCP1XX [2288892]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,3 MEDLINE links, 28 protein links, 1 nucleotide neighbor, or 1 genome link)

L35561

Bacteriophage phi-105 ORFs 1-3
gil532218|gb|L35561|PH5ORFHTR [532218]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 3 protein links)

D49712

Bacillus licheniformis DNA for ORFs, xpaL2 homologous protein and xpaL1 homologous protein, complete and partial cds
gil1514423|dbj|D49712|D49712 [1514423]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, or 4 protein links)

X90511

Lactobacillus bacteriophage phig1e DNA for Rorf162, Holin, Lysin, and Rorf175 genes
gil1926386|emblX90511|LBPHIHOL [1926386]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein links, or 1 nucleotide neighbor)

X98106

Lactobacillus bacteriophage phig1e complete genomic DNA
gil1926320|emblX98106|LBPHIG1E [1926320]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

link, 50 protein links, or 4 nucleotide neighbors)²⁵⁹

U72397

Bacteriophage 80 alpha holin and amidase genes, complete cds
gil1763241|gb|U72397|B8U72397 [1763241]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 protein links, or 2 nucleotide neighbors)

U38906

Bacteriophage rlt integrase, repressor protein (rro), dUTPase, holin and lysin genes, complete cds
gil1353517|gb|U38906|BRU38906 [1353517]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,2 MEDLINE links, 50 protein links, or 3 nucleotide neighbors)

X91149

Bacteriophage phi-C31 DNA cos region
gil1107473|embl|X91149|APHIC31C [1107473]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 6 protein links, or 1 nucleotide neighbor)

U24159

Bacteriophage HP1 strain HP1c1, complete genome
gil1046235|gb|U24159|BHU24159 [1046235]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,6 MEDLINE links, 41 protein links, 8 nucleotide neighbors, or 1 genome link)

Z26590

Bacteriophage mv4 lysA and lysB genes
gil410500|embl|Z26590|MV4LYSAB [410500]
(View GenBank report,FASTA report,ASN.1 report,Graphical view, or 4 protein links)

Z70177

B.subtilis DNA (28 kb PBSX/skin element region)
gil1225934|embl|Z70177|BSPBSXSE [1225934]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,32 protein links, or 4 nucleotide neighbors)

Z36941

260

B.subtilis defective prophage PBSX xhlA, xhlB, and xylA genes
gil535793|embl|Z36941|BSPBSXXHL [535793]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,4 protein
links, or 5 nucleotide neighbors)

X89234

L.innocua DNA for phagelysin and holin gene
gil1134844|embl|X89234|LICPLYHOL [1134844]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 2 protein links, or 4 nucleotide neighbors)

X85010

Bacteriophage A511 ply511 gene
gil853748|embl|X85010|BPA511PLY [853748]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 3 protein links, or 1 nucleotide neighbor)

X85009

Bacteriophage A500 hol500 and ply500 genes
gil853744|embl|X85009|BPA500PLY [853744]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 3 protein links, or 4 nucleotide neighbors)

X85008

Bacteriophage A118 hol118 and ply118 genes
gil853740|embl|X85008|BPA118PLY [853740]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 3 protein links, or 1 nucleotide neighbor)

L34781

Bacteriophage phi 11 holin homologue (ORF3) gene, complete cds and
peptidoglycan hydrolase (lytA) gene, partial cds
gil511838|gb|L34781|BPHHOLIN [511838]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE
link, 4 protein links, or 2 nucleotide neighbors)

U11698

Serratia marcescens SM6 extracellular secretory protein (nucE), putative
phage lysozyme (nucD), and transcriptional activator (nucC) genes,
complete cds
gil509550|gb|U11698|SMU11698 [509550]
(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE

link, 3 protein links, or 1 nucleotide neighbor)

U31763

Serratia marcescens phage-holin analog protein (regA), putative phage lysozyme (regB), and transcriptional activator (regC) genes, complete cds

gi1965068|gb|U31763|SMU31763 [965068]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 3 protein links, or 1 nucleotide neighbor)

X87674

Bacteriophage P1 lydA & lydB genes

gi1974763|emb|X87674|BACP1LYD [974763]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 2 protein links, or 2 nucleotide neighbors)

L48605

Bacteriophage c2 complete genome

gi1146276|gb|L48605|C2PVC [1146276]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 3 MEDLINE links, 39 protein links, 3 nucleotide neighbors, or 1 genome link)

L33769

Bacteriophage bIL67 DNA polymerase subunit (ORF3-5), essential recombination protein (ORF13), lysin (ORF24), minor tail protein (ORF31), terminase subunit (ORF32), holin (ORF37), unknown protein (ORF 1-2, 6-12, 14-23, 25-30, 33-36), complete genome

gi522252|gb|L33769|L67CG [522252]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 37 protein links, 2 nucleotide neighbors, or 1 genome link)

L31348

Bacteriophage Tuc2009 integrase (int) gene, complete cds; lysin (lys) gene, 3' end

gi508612|gb|L31348|TU2INT [508612]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 3 protein links, or 3 nucleotide neighbors)

L31364

Bacteriophage Tuc2009 holin (S) gene, complete cds; lysin (lys) gene, complete cds

gi496281|gb|L31364|TU2SLYS [496281]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

L31366

Bacteriophage Tuc2009 structural protein (mp2) gene, complete cds
gi496278|gb|L31366|TU2MP2A [496278]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

L31365

Bacteriophage Tuc2009 structural protein (mp1) gene, complete cds
gi496276|gb|L31365|TU2MP1A [496276]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, or 1 protein link)

U04309

Bacteriophage phi-LC3 putative holin (lysA) gene and putative murein
hydrolase (lysB) gene, complete cds

gi530796|gb|U04309|BPU04309 [530796]

(View GenBank report,FASTA report,ASN.1 report,Graphical view,1 MEDLINE link, 2 protein links, or 1 nucleotide neighbor)

Table 14

NCBI *Entrez* Nucleotide QUERY

Key word: bacteriophage and kil

5 citations found (all selected)

AF034975

Bacteriophage H-19B essential recombination function protein (erf), kil protein (kil), regulatory protein cIII (cIII), protein gp17 (17), N protein (N), cI protein (cI), cro protein (cro), cII protein (cII), O protein (O), P protein (P), ren protein (ren), Roi (roi), Q protein (Q), Shiga-like toxin A (slt-IA) and B (slt-IB) subunits, and putative holin protein (S) genes, complete cds
gil2668751|gb|AF034975| [2668751]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 20 protein links, or 30 nucleotide neighbors)

X15637

Bacteriophage P22 P(L) operon encompassing ral, 17, kil and arf genes
gil15646|emb|X15637|POP22PL [15646]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 7 protein links, or 2 nucleotide neighbors)

J02459

Bacteriophage lambda, complete genome
gil215104|gb|J02459|LAMCG [215104]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 87 MEDLINE links, 67 protein links, 190 nucleotide neighbors, or 1 genome link)

M64097

Bacteriophage Mu left end
gil215543|gb|M64097|PMULEFTEN [215543]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 2 MEDLINE links, 39 protein links, or 15 nucleotide neighbors)

M18902

Bacteriophage D108 kil gene encoding a replication protein, 3' end; and containing three ORFs, complete cds
gil166191|gb|M18902|D18KIL [166191]

(View GenBank report, FASTA report, ASN.1 report, Graphical view, 1 MEDLINE link, 1 protein link, or 3 nucleotide neighbors)

Table 15

U77328	V01282	U11787	U93688	A47599	D21131	U76864	U38428
AF151117	AF121672	U11786	U93687	A47598	D30690	U76863	U66665
AF151218	AF072726	U11785	AJ224764	A47597	D14711	U76862	U66664
AF146368	AF115379	U11784	AF064774	A47596	D90119	U76861	U66663
AF144661	AF034153	U11783	AF064773	A47595	D00730	U76860	X87104
AF132117	AF029244	U11782	Y14370	A47594	D83357	U76859	X87105
Y15477	U67965	U11781	AF065394	A44534	D83356	U76858	X89233
Y09928	U96610	U11780	AF062376	A44533	D83355	U76857	M28521
Y09594	U96609	U11779	AF062375	A44529	D83354	U76855	U54636
AF134905	U73027	U11778	AF062374	A44528	D83353	U76854	U46541
AB019536	U73026	U11777	AF062373	A44527	D12572	U76853	L14017
AJ237696	U73025	U11776	AB007500	A44526	D86727	U76852	U60589
AF106851	AF068904	U11775	Y09924	A44525	D86240	U76851	X48003
AF106850	U60050	U11774	U63529	A39696	D67075	U76850	M37889
AF106849	D10907	U11773	AF033191	AF001783	D67074	U76849	V01281
M26321	D10906	AF053772	Y15856	AF001782	U97062	U76848	X97985
AF060191	AF053140	AF053771	AB000439	L77194	U96620	U76847	X00127
AF060190	AB013298	AF029731	AF041467	AF003593	U96619	Y09929	X03286
AF060189	Y16431	AF027155	Y14051	AF003592	Z84573	Y09570	X62282
AF060188	AF076684	AF024571	U82085	X73889	AB001896	X95848	X01645
AF060187	AF076683	U87144	AF026122	X74219	Y07645	Y09428	X16471
AF060186	Y13225	AF086644	AF026121	Y10419	U92441	S76611	X52734
AF060185	AF094826	AJ223781	AF026120	M63177	U91741	S76213	X13290
AF060184	AJ223480	AF076030	AB009635	E08773	U29454	S75707	X66088
AF036324	AF093548	AF044951	AB006796	E07163	U29478	S75706	Z30588
AF036323	AJ005352	AF044906	U39769	E07162	U77374	S75705	X16457
AF053568	AF051916	AF044905	D00184	E07161	L42945	S76270	X00342
AJ132841	Y09927	AF044904	X56628	E07160	U38429	S72497	V01287
Y13766	AF051917	AF044903	AF033018	E07159	U81980	S72488	X61307
AF101234	S77058	AF044902	AF034076	E07158	X55185	S74031	Y00356
AJ133520	S65052	AF044901	D82063	E07157	V01278	S67449	X06603
AJ133495	AF009671	AF044900	D76414	E07156	U31979	U75367	Z93205
AJ132803	U81973	AF044899	U57060	E07155	X91786	U75368	X64172
AB016487	U77308	AF044898	D89066	E03836	U36912	U31175	X72700
AB016431	U20869	AF044897	U85095	E03835	U36911	X53096	X60827
AB015981	U89396	AF044075	U85097	E03526	U36910	X53951	X64389
AB015195	U94706	AF044074	U85096	E02873	U64885	X53952	X62288
AF107307	U41072	AF044073	D42078	E01690	U76872	X03408	X55798
AF079518	U52961	AF044072	AF015929	E00876	U76871	U50629	X58434
AJ223806	U21636	AF044071	D10369	E00203	U76870	U38656	X06627
Y18018	U65000	AF044070	A48955	D83951	U76869	U58139	X12831
Y17795	U48826	AF044069	A48501	D17366	U76868	A31894	X07371
AJ005647	U20503	AF044068	A48500	D42144	U76867	L42943	X02529
AJ005646	U11789	AF044067	A48499	D42143	U76866	U51474	Y00688
AJ005645	U11788	AF044066	A47600	D10489	U76865	U50077	X04121
X59477	X54338	A12915	U51133	M63176	M10500	L01055	M63917
X59478	X51661	A12913	U51132	L11998	M10499	M83994	M58515
X63598	X05815	A12906	X02588	L05004	AH000934	J03947	L10909
X52593	X15574	A12905	X61716	L42764	M10498	J03479	M15067

X76490	Y07536	A12904	X61719	M32103	M10497	M64724	M92376
X81586	X02166	A12903	X61718	U10927	M18264	M14372	M62650
X72014	Z49245	A12902	X67743	AH003057	J01786	M14371	M32312
X72013	X16298	A12901	X67742	M73535	M33833	M14374	M20393
X71437	Z18852	A12900	X67741	M73536	M32470	M15215	M90536
X62992	X68417	A12899	X67740	U20782	M20270	M36694	M21854
X52594	X68425	A12898	X67738	L37598	J03323	M37915	M36771
X14827	X17679	A12897	U02910	L37597	M33479	M12715	L14020
X13404	X63072	A12896	AH003349	L36472	M94061	J04151	M81736
X17301	X02872	A09523	M11118	L25288	M37888	L22566	U11702
X17688	V01277	A04518	M18086	L25893	M76714	L13379	L19300
X03097	X52543	A04517	U19459	K02687	M17123	L13378	L25372
Z16422	A19943	A04512	U35773	L23109	M97169	L13377	L22565
Z33409	A19942	L41499	U26702	L07778	M81346	L13376	M58516
Z33408	A19941	U19770	U21221	M90056	M90693	L13375	U06462
Z33407	A19940	X53818	U36379	J02615	M25257	L13374	L19298
Z33406	A19939	M20129	U06451	M18970	M25256	M17348	M80252
Z33405	A19938	L43098	U35036	K02985	M25255	M17357	L11530
Z33404	A19937	L43082	U20794	M21136	M25254	M17347	
X75439	A19936	X03216	L25426	M10501	M25253	M28364	
X62587	A17958	X70648	M86227	AH000935	M25252	M21319	

Table 16

Phage 44AHJD complete genome sequence. 16668 nucleotides.

```

1      tccatttctt tactaaactt aaaaatgctg tgcaacaact taaccaactt atctaacta ttacatattc
71     atcaaataca aaattttatgt atctattgac ttttattcaa aattatgatt tcaacatata ataaaaattaa
141    tttacttatt taatatattct atgataataa tagttataaa atatttgag gtgtataaat gacagaattt
211    gatgaaatcg taaaaccaga cgacaaagaa gaaacttcag aatcaactga agaaaattta gaatcaactg
281    aagaaacttc agaatacaact gaagaatcaa ctgaagaatc aactgaagaa tcaactgaag ataaaaacagt
351    agaaacaact gaagaagaaa atgaaaacaa attagaacct actacaacag atgaagatga ttcgaaattt
421    gaccctgttg tattagaaca acgtattgct tcattagaac aacaagtgc tactttttta tcttcacaaa
491    tgcaacaacc acaacaagta caacaaacac aatcagatgt aacagaatca acaaaagaag ataactgacta
561    ttcagatgaa gaactagttg ataagttaga tttagattag gaggaattta aacatgtatg agggaaacaa
631    catgcgttct atgatgggta catcatatga agattcaaga ttaataaacc gaacagaatt aaatgaaaac
701    atgtcaattg atacaaataa aagtgaagat agttatgggt tacaatttca ttcactttca aaacaatcat
771    ttacaggtga cgttgaggag gaataataaa ttatggcaca acaatctaca aaaaatgaaa tgcacttttt
841    agtagcaaaag tcagctaaat cagcgttaca agattttaat catgattatt caaaatcttg gacatttggc
911    gacaaatggg ataattcaaa tacaatgttc gaaacatttg taaataaata tttattccct aagattaatg
981    agactttatt aatcgatatt gcattaggta atcgttttaa ttggttagct aaagagcaag attttattgg
1051   acaatatagt gaagaatcag tgattatgga cacagtacca attaacatgg acttatctaa aatgaggaaa
1121   ttaatgttga aacgtaatta tccacgtatg gcaactaagt tatatggtta cgggaatttg aagaaacaaa
1191   aattcacatt aaacaacaat gatcacggtt caattttcca aacattagca gacgcaacta attacgcttt
1261   aggtgtatc aaaaagaaaa tttctgatat taatgtatta gaagaaaaag aaatgcgtgc aatgttagtt
1331   gattactcat tgaatcaatt atccgaacaa aatgtacgta aagcaacatc aaaagaagat tttagcaagca
1401   aagtttttga agcaatccta aacttaccaa acaacagtgc taatatataat gaagtacatc gtgcatcagg
1471   tgggtcaatt ggacaatata caactgtatc aaaattaaaa gatattgtga ttttaacacac agattcatta
1541   aaatcttate ttttagatac taagatttga aacacattcc agattgcagg cattgatttc acagatcacg
1611   ttattagttt tgacgactta ggtggcgtgt ttaaagtaac aaaagaattt aagttacaaa accaagattc
1681   aattgacttt ttacgtgcgt atggagatta tcaatcacaa ttaggagata caattccagt tgggtgctga
1751   ttactattag atgtatctaa acttaagag ttacttgcca acgttgaaaga aattaaacca aaatcagatt
1821   tatatgcgtt tattttggat attaatctaa ttaaatataa acgttacaca aaaggtatgt taaaaccacc
1891   attccataac cctgaatttg atgaagttac aactggatt cattactatt catttaaacg cattagtcca
1961   ttctttaata aaattttaat tactgaccaa gatgtaaatc caaaaccaga ggaagaatta caagaataaa
2031   agggacgtta aatatgaaca acgataaaaag aggtttaaac gttgagttat caaaggaatc cagcaaaaaga
2101   gttgttgaa atcgcaacag atttaaacgt cttatgttta atcgttattt ggaattttta ccgctactaa
2171   tcaactatac caatcgtgat acggttggtg tagattttat tcagttagaa tcagctttta gacaaaacat
2241   taatgtagtt gttggtgaag ctagaataaa gcaaatatag attcttggtt atgtaataaa caacttattt
2311   aatcaagcac caaatttttc atcaaacctt aatttccaat ttcaaaaacg attaaactaa gaagatatat
2381   attttattgt acctgactat ttaatacctg atgattgtct acaaatctat aagctatag ataactgtat
2451   gagtggtaac tttgtgtcca tgcaaaaata accaattcaa tataatagtg atatagaatc atatagaacat
2521   tatactgatg aattagcaga agttgcttta tctcgctttt cttaaatcat gcaagcaaaa tttagcaaga
2591   tatttaaatc agaaattatg gacgagtcac tcaatcaact tgtgtccgaa atatataacg gtgcaccatt
2661   tgttaaaatg tcacctatgt ttaatgcaga tgacgatatc attgatttaa caagtaaatg cgtaatccca
2731   gcattaactg aaatgaaacg ggaatatcaa aacaaaatta gtgaattaag taactattta ggcattaatt
2801   cattagccgt tgataaagaa agcgtgtgtt cagacgaaga ggcaaaaagt aatcggtggat ttaccacatc
2871   aaacagtaat atctatttaa aaggtcgtga accaattacg tttttatcaa agcgttatgg tttagattat
2941   aaacgctatt acgatgatga aacaacgtct aaaaattcaa tggtagacac actttttaa gatgaagca
3011   gtgataataa tggctagata cacaatgact ttatcagatt tcattaaatc agaattgatt aaaaagggtt
3081   tcaatgaatt tgtaaatgat aataaattaa cgttttatga tgatgaattt caattcatgc aaaaaatgct
3151   gaagttcgac aaagacgttt tagctatcgt taatgaaaaa gtattttaaag gtttttcatt gaaagatgaa
3221   ttatcagatt tactttttta aaaatcattt acgattcatt ttttagatag agaaatcaac agacaaacag
3291   ttgaagcatt tggcatgcaa gtgattactg tatgtattac acatgaggat tatttaaatg tggttttatc
3361   atcaagtga gttgaaaaat acttacaatc acaaggcttc acagaacaca atgaagatac aacaaatgaa
3431   actgatgaaa catcgaaatc aaatgctaca tctttagaca attcaactgg catgactgca aacagaaacg
3501   cttatgtgtc attaccacaa agtgaggtta acattgatgt tgataatata acgttacgat tccgtgataa
3571   taatacgaat gataacggta aaactgtgaa taaatcgagt aacgaaagta atcaaaacgc aaaacgtaat
3641   caaaatcaaa aaggtaatgc aaaaggtaca caattcacta agcagttatt aattgataat attgataaag
3711   cgtacgattt aagaaagaaa attttaaatg aatttgataa aaaatgtttt ttacaaattt ggtagaggtg
3781   gttaaataat ggcataataa gaaaacgatt ttaaatattt tgatgacatt cgtccatttt tagacgaaat
3851   ttataaaacg agagaacggt atacacggtt ttacgatgat agagcagatt ataatactaa ttcaaatca
3921   tattatgatt atatttcaag attatcaaaa ctaattgaag tattagcacg tctgatttgg gactatgaca
3991   atgaattaaa aaaacgtttc aaaaattggg acgacttaat gaaagcattt ccagagcaag cgaagactt
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4481   tgggtgaaatg aaaatctggt tacatcacga tgggtgtgca aaactgttac aagtcgcata taaagataat
4551   tatgtattag atttgaaga ggctaaaggt tttaacgatt ataccacaca gtcactttta aacaaacaca
4621   catttacacc gtaattgat gaagcaaatg acaaatcat ttttaagatt ggtgacggaa caatacaggt
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16661 ggtggggt

Table 17

Phage 44AHJD ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	44AHJDORF001	-1	10342..12627	761	DNA polymerase;
2	44AHJDORF002	3	3789..5732	647	Teichoic acid; Staph;
3	44AHJDORF003	2	6626..8389	587	Tail;
4	44AHJDORF004	1	8764..10227	487	Serine protease motif;
5	44AHJDORF005	-1	12643..13890	415	
6	44AHJDORF006	2	803..2029	408	
7	44AHJDORF007	1	2044..3027	327	Upper collar;
8	44AHJDORF008	2	3020..3775	251	Lower collar;
9	44AHJDORF009	2	5744..6496	250	Amidase; Staph;
10	44AHJDORF010	-2	13938..14420	160	
11	44AHJDORF012	3	8391..8813	140	Holin;
12	44AHJDORF013	-2	14586..14996	136	
13	44AHJDORF113	1	199..600	133	
14	44AHJDORF011	-2	15225..15593	122	
15	44AHJDORF114	-2	15870..16172	100	
16	44AHJDORF014	3	6243..6521	92	
17	44AHJDORF015	1	15403..15645	80	
18	44AHJDORF016	-1	15616..15852	78	
19	44AHJDORF017	-2	10536..10757	73	
20	44AHJDORF018	-1	886..1098	70	
21	44AHJDORF019	-2	9630..9836	68	
22	44AHJDORF121	-1	16165..16362	65	
23	44AHJDORF020	2	13865..14053	62	
24	44AHJDORF123	2	614..796	60	
25	44AHJDORF021	-2	5634..5816	60	
26	44AHJDORF023	-2	6315..6494	59	
27	44AHJDORF024	1	14275..14451	58	
28	44AHJDORF025	-3	14999..15175	58	
29	44AHJDORF026	-3	14426..14593	55	
30	44AHJDORF027	1	12916..13080	54	
31	44AHJDORF029	-1	15019..15183	54	
32	44AHJDORF028	-3	9071..9235	54	
33	44AHJDORF030	3	14487..14648	53	
34	44AHJDORF031	2	11039..11191	50	
35	44AHJDORF135	3	693..842	49	
36	44AHJDORF033	-1	3646..3795	49	
37	44AHJDORF032	-2	9306..9455	49	
38	44AHJDORF034	-3	14000..14146	48	
39	44AHJDORF035	-3	13811..13957	48	
40	44AHJDORF036	-3	10019..10165	48	
41	44AHJDORF022	-3	8468..8611	47	
42	44AHJDORF037	1	14788..14931	47	
43	44AHJDORF038	-2	3528..3671	47	
44	44AHJDORF039	3	1743..1883	46	
45	44AHJDORF040	2	9740..9877	45	
46	44AHJDORF041	2	15836..15973	45	
47	44AHJDORF042	-1	5014..5151	45	
48	44AHJDORF043	-1	4402..4539	45	
49	44AHJDORF044	-2	12783..12917	44	
50	44AHJDORF149	-2	639..770	43	
51	44AHJDORF046	1	4891..5019	42	
52	44AHJDORF047	1	11911..12039	42	
53	44AHJDORF045	2	10655..10783	42	
54	44AHJDORF048	-3	15212..15340	42	
55	44AHJDORF049	3	5784..5909	41	
56	44AHJDORF050	3	13158..13283	41	
57	44AHJDORF051	-2	10944..11066	40	
58	44AHJDORF052	-3	14216..14338	40	
59	44AHJDORF053	3	3348..3467	39	
60	44AHJDORF054	3	7551..7670	39	
61	44AHJDORF055	3	15705..15821	38	
62	44AHJDORF056	1	5512..5625	37	
63	44AHJDORF057	2	10121..10231	36	
64	44AHJDORF058	3	10767..10877	36	

271

65	44AHJDORF164	-1	592..702	36	
66	44AHJDORF059	-2	8250..8360	36	
67	44AHJDORF060	-2	6147..6257	36	
68	44AHJDORF061	2	15551..15658	35	
69	44AHJDORF062	1	4285..4389	34	
70	44AHJDORF063	-3	9383..9487	34	
71	44AHJDORF065	1	5029..5130	33	
72	44AHJDORF064	2	2609..2710	33	
73	44AHJDORF066	-2	10380..10481	33	

Table 18

Predicted amino acid sequences

44AHJDORF001

12627 atgggattactagaatgcacatgcataaacatgaacgtcgaatgattttatactgggatagaaacattagcgtacaat
 1 M G L L E C M Q Y H K H E R R M I L Y W D I E T L A Y N
 12543 aaagttaacggacgaaaaaaacaaacaaatataaaaacgttacttattctgtagcaattggttggttaaggttatgaaatt
 29 K V N G R K K P T K Y K N V T Y S V A I G W F N G Y E I
 12459 gatgttgaagtatttccgagtttcgaatctttttatgcacgattttatacgtatgtgaaaagacgtgatacaatcacaaaatca
 57 D V E V F P S F E S F Y D A F Y T Y V K R R D T I T K S
 12375 aaaacagatattatcatgattgcacataactgtaataaacatgcataatcatttttacttaagacaccatgcgttattttgat
 85 K T D I I M I A H N C N K Y D N H F L L K D T M R Y F D
 12291 aatattacacgcgaaaaatataatatttaaaatctgcagaagaaaatgaacacacattaaaaatgaaagaggctactattttagcc
 113 N I T R E N I Y L K S A E E N E H T L K M K E A T I L A
 12207 aaaaatcaaaatgtaatttttagaaaaacgtgttaaatcttcaatcaatttagatttaacaatgttttttaaatggttttaaat
 141 K N Q N V I L E K R V K S S I N L D L T M F L N G F K F
 12123 aatattattgataactttatgaaaacaaatcatcaattgcaacattaggttaagaaattacttgatggtggttatttaacagaa
 169 N I I D N F M K T N T S I A T L G K K L L D G G Y L T E
 12039 tcacaacttaaaacagattttaattatacagatttttgataaagataatgatgatgatgtaggaacccatgactatgctgtg
 197 S Q L K T D F N Y T I F D K D N D M N D S E A Y D Y A
 11955 aaatgttttgcaaaactcacacctgaacaaactcacacattacataatgacgtgatttatattaggtatgtgccatattcattat
 225 K C F A K L T P E Q L T Y I H N D V I I L G M C H I H Y
 11871 agtgatatatttccaaattttgactatacaaaatcaacatttttctgaatattatggaatcttacttgaataatgaaatgaca
 253 S D I F P N F D Y N K L T F S L N I M E S Y L N N E M T
 11787 cgttttcagttactcaaccaatatacaagattataaaatcttatacacattatcatttccatgatgatgaatttttatgactat
 281 R F Q L L N Q Y Q D I K I S Y T H Y H F H D M N F Y D Y
 11703 attaaatcattctatcgtggtggttttaaatatgtataacacaaataacataaaactaattgatgagcctgtttttctatt
 309 I K S F Y R G G L N M Y N T K Y I N K L I D E P C F S I
 11619 gacatcaattcaggttatccttatgtgatgatcatgaaaaaattccaacatggttatacttttcaaacactattcagaacca
 337 D I N S S Y P Y V M Y H E K I P T W L Y F Y E H Y S E P
 11535 acgttaactccctacttttttagatgatgacaattattttcattataaagattgataaagatgtatttaacgatgatttatta
 365 T L I P T F L D D D N Y F S L Y K I D K D V F N D D L L
 11451 attaaaaataaacacgtgtattacgtcaaatgattgttaaaataactataaataatgataatgattacgttaatatcaatacaaat
 393 I K I K S R V L R Q M I V K Y Y N N D N D Y V N I N T N
 11367 acattaagaatgattcaagacattacgggtattgattgcacatcatcgtgttaattcgtttgttatatgatgatgtgaatcac
 421 T L R M I Q D I T G I D C M H I R V N S F V I Y E C E Y
 11283 tttcatgcacgtgatattatttttcaaaactattttattaaacacaaaggttaagttaaaaacaaatcaatatgacatcacct
 449 F H A R D I I F Q R N Y F I K T Q G K L K N K I N M T S P
 11199 tacgactatcacattactgatgatcatcaacgaacaccataactcaaatgaggaggttatgttatctaaagtcgttttaaatgga
 477 Y D Y H I T D D I N E H P Y S N E E V M L S K V L N G
 11115 ttatatggcatacctgcattacgttcacatttttaacttattccgttttagatgataaactgaactatacaatatcattaacggt
 505 L Y G I P A L R S H F N L F R L D D N N E L Y N I N G
 11031 tacaaaaacactgaacgtaatatattattctctacattttgtcacatcagttcattgtataacttattggttccctttccatcac
 533 Y K N T E R N I L F S T F V T S R S L Y N L L V P F Q Y
 10947 ttaacggaaagtgaatttgacgacaattttatttattgacgatactgatagtttgatatgaaatccgttggttaaaccccttattg
 561 L T E S E I D D N F I Y C D T D S L Y M K S V V K P L L
 10863 aaccccgagtttattcgaccgatagccttaggttaattgggatattgaaaacgaacagatagataagatggttgactgaatcat
 589 N P S L F D P I A L G K W D I E N E Q I D K M F V L N H
 10779 aagaaatgatcatatgaagtgaatggaagattaaaattgcttctgctggtataccgaaaaacgcctttgatacaagcgtcgat
 617 K K Y A Y E V N G K I K I A S A G I P K N A F D T V D
 10695 tttgaaacctttgtacgtgaacaattctttgacggtgccattattgaaaacataaaaagtatctataatgagcaaggtacaata
 645 F E T F V R E Q F F D G A I I E N N K S I Y N E Q G T I
 10611 tcgatatatccgtctaaaactgaaattgtatgtggaatgtatatgatgaatattttactgatgaacttaatatgaaacgtgaa
 673 S I Y P S K T E I V C G N V Y D E Y F T D E L N M K R E
 10527 tttatattaaaaagacgctagagaaaaatttcgaccatagtcattttgatgatattctttatattgaaagdtgacatcggttcatt
 701 P I L K D A R E N F D H S Q F D D I L Y I E S D I G S F
 10443 tcacttaacgacttatttccagttgaacgttcagttacataaacaattctgatttgcatattataaacgtgaacatgatgaata
 729 S L N D L F P V E R S V H N K S D L H I L K R E H D E I
 10359 aaaaaaggcaactgttaa 10342
 757 K K G N C *

44AHJDORF002

3789 atggcatataatgaaaacgatttttaaatattttgatgacattcgtccatttttagacgaaatttataaaacgagagaacggttat
 1 M A Y N E N D F K Y F D D I R P F L D E I Y K T R E R Y
 3873 acaccggttttacgatgatagacagattataataactaattcaaaatcatattatgattatatttcaagattatcaaaactaatt
 29 T P F Y D D R A D Y N T N S K S Y Y D Y I S R L S K I
 3957 gaagtatttagcagctgattttgggactatgacaatgaattaaaaaacggtttcaaaaattgggacgacttaagtgaagacttt
 57 E V L A R R I W D Y D N E L K K R F K N W D D L M K A F
 4041 ccagagcaagcgaaagacttatttagaggttggttaaacgacggtacgattgacagattattcatgacgagtttaaaaaatatt
 85 P E Q A K D L F R G W L N D G T I D S I I H D E P K K Y
 4125 agcgcaggatttaacatcggttcttttatttaagtttactgaaatgaaacaaatgaatgacttttaaatcagaagtttaagac
 113 S A G L T S A F A L F K V T E M K Q M N D F K S E V K D
 4209 ttaattaaagatattgaccggttctgtaattgggtttgaattaaatgagcttgaaacaaagtttgatgaggttcttggtggtatt

275

803 atggcacaacaatctacaaaaatgaaactgcacttttagtagcaaagtcagctaaatcagcgttacaaagattttaatcatgat
1 M A Q Q S T K N E T A L L V A K S A K S A L Q D F N H D
887 tattcaaaatcttgacatttggcgacaaatgggataattcaaatcaaatgttcgaaacatttggtaataaatatttattccct
29 Y S K S W T F G D K W D N S N T M F E T F V N K Y L F P
971 aagattaatgagacttttataatcgatattgcattaggttaactgttttaattgggttagctaaagagcaagattttattggacaa
57 K I N E T L L I D I A L G N R F N W L A K E Q D F I G Q
1055 tatagtgaagaatacgtgattatggacacagtagcaatcaatcaggttattctaaaaatgaggaattatgttgaaacgtaat
85 Y S E E Y V I M D T V P I N M D L S K N E E L M L K R N
1139 tatccacgtatggcaactaagttatatgttgtaacggaattgtgaagaacaaaaattcacattaaacaacatgatacagtttc
113 Y P R M A T K L Y G N G I V K K Q K F T L N N N D T R F
1223 aatttccaaacattagcagacgcaactaattacgtcttaggtgtatacaaaaagaaaatttctgatattaatgtattagaagaa
141 N F Q T L A D A T N Y A L G V Y K K K I S D I N V L E E
1307 aaagaaatgcgtgcaatgttagttgattactcattgtaactcaattatccgaaacaaatgtacgtaaagcaacatcaaaagaagat
169 K E M R A M L V D Y S L N Q L S E T N V R K A T S K E D
1391 tttagcaagcaagtttttgaaagcaatcctaaactcaaaacaacagtgctaaatataatgaagtacatcggtgcacaggtggg
197 L A S K V P E A I L N L G N N S A K Y N E V H R A S G G
1475 gcaattggacaatatataactgtatcaaaatataagatattgtgattttaacaacagattcattaaaaatcttatttttagat
225 A I G Q Y T T V S K L K D I V I L T T D S L K S Y L L D
1559 actaagattgcaaacacattccagattgcaggcattgtattcacagatcaggttatttagtttgacagattaggtggcgtgtt
253 T K I A N T F Q I A G I D F T D H V I S F D D L G G V F
1643 aaagtaacaaagaatttaagttacaaaaccaagattcaattgactttttacgtgcgtatggagattatcaatcacattagga
281 K V T K E F K L Q N Q D S I D F L R A Y G D Y Q S Q L G
1727 gatacaattccagttgggtgtgtatttactttatgattatcctaaacttaaaagagtttactggcaacgttggaataataaacca
309 D T I P V G A V F T Y D V S K L K E F T G N V E E I K P
1811 aaatcagatttatatgcgttttattttggatattaattcaattataaaacgttacacaaaagggtattgttaaaaccaccattc
337 K S D L Y A P I L D I N S I K Y K R Y T K G M L K E D I
1895 cataaccctgaatttgatgaagttacacactggattcattactattcatttaaaagccatttagtccattctttaataaaatttta
365 H N P E F D E V T H W I H Y Y S P K A I S P F P N K I L
1979 attactgaccaagatgtaaatccaaaaccagaggaagaattacaagaataa 2029
393 I T D Q D V N P K P E E E L Q E *

44AHJDORF007

2044 atgaacaacgataaaaagaggtttaaaacgttgagttatcaaaggaatcagcaaaaagagttgtgaacatcgcaacagatttaaa
1 M N . N D K R G L N V E L S K E I S K R V V E H R N R F K
2128 cgtcttatgtttaatcgttatttggaaattttaccgctactaatcaactataccaatcgtgatacgggttggtatagattttatt
29 R L M F N R Y L E F L P L L I N Y T N R D T V G I D F I
2212 cagttagaatcagctttaagacaaaacatttaagtgagttgttggaagctagaataagcaaatatttctgttattgtatga
57 Q L E S A L R Q N I N V V V G E A R N K Q I M I L G Y V
2296 aataacacttactttaatcaagcaccacaaatttttcatcaaaactttaatttccaatttcaaaaacgatttaactaaagaagata
85 N V T F N Q A P N F S I N N F Q F Q K R L T K E D I
2380 tattttattgtacctgactatttaatacctgatgattgtctcaaaattcataagctatatgataactgtatgagtggttaacttt
113 Y F I V P D Y L I P D D C L Q I H K L Y D N C M S G N F
2464 gttgtcatgcaaaaataaaccattcaatataatgtagatagaataatagaaacattatagaaactgaggaatttagcaggaattgct
141 V V M Q N K P I Q Y N A S D I E I I E H Y T D E L A E V A
2548 ttatctcgctttttttaaattcatgcaagcaaaaatttagcaagatatttaaatcagaatattgacgagtcattcaatcaactt
169 L S R F S L I M Q A K F S K I F K S E I N D E S I N Q E
2632 ggtgccgaaatataacggtgcaccatttgttaaaattgtcacctatgtttaatgcagatgacgatattcattgatttacaag
197 V S E I Y N G A P F V K M S P M F N A D D D I I D L T S
2716 aatagcgttaatccagcatttaactgaaatgaaacgggaatatacaaaaacaaatttagtgaattaagtaactatttaggcattaat
225 N S V I P A L T E M K R E Y Q N K I S E L S N Y I N
2800 tcattagccgttgataaagaaagcgtgtttcagacgaagaggcaaaaagtaactcgtggatttaccacatcaaacagtaatatc
253 S L A V D K E S G V S D E E A K S N R G F T T S N S N I
2884 tatttaaaaggtcgtgaaccaattacgttttttaacagcgtttatggttttagatattaaacgttattacgatgataaacaag
281 Y L K G R E P I T F L S K R Y G L D I K P Y Y D D E T T
2968 tctaaaaatataatggttagacacactttttaagatgaaagcagtgatataaatggctag 3027
309 S K I S M V D T L F K D E S S D I N G *

44AHJDORF008

3020 atggctagatacacaaatgactttatcagatttcttaaatcagaattgattaaaaaagggtttcaatgaatttgtaaatgataat
1 M A R Y T M T L Y D F I K S E L I K K G F N E F V N D N
3104 aaattaacgttttatgatgatgaatttcaattcatgcaaaaaatgctgaagttcgacaaagcgttttagctatcgtaaatgaa
29 K L T F Y D D E F Q F M Q K M L K F D K D V L A I V N E
3188 aaagtatttaaggtttttcattgaaagatgaattacagatttacttttaaaaaatcatttacgattcatttttttagataga
57 K V F K G F S L K D E L S D L L F K K S F T I H F L D R
3272 gaaatcaacagacaaacagttgaagcatttggcatgcaagtgtattactgtatgtattacacatgaggattatttaaatgtggtt
85 E I N R Q T V E A F G M Q V I T V C I T H E D Y L N V V
3356 tattcatcaagtgaaagtgaaaaacttacaacacagaaggttcacagaacacaaatgaagatacaacaagtaacacagtgaa
113 Y S S S E V E K Y L Q S Q G F T E H N E D T T S N T D E
3440 acatcgaatcaaaatgctacatcttttagacaattcaactggcagtgactgcaaacagaaacgcttatgtgtcattaccacaag
141 T S N Q N A T S L D N S T G M T A N R N A Y V S L P Q S
3524 gaggttaacattgatgttgataatacaacggttacgattcgtgataataatcagattgataacggtaaaactgtgaataaatcg.
169 E V N I D V D N T T L R F A D N N T I D N G K T V M L K S
3608 agtaacgaaagtaatacaaacgcaaacgtaatacaaaatgaaaggttaacgaaaggttacacaattcactaagcagatttta
197 S N E S N Q N A K R N Q N Q K G N A K G T Q F T K Y L
3692 attgataatattgataaagcgtacgatttaagaagaaaaattttaaatgaatttgataaaaaatggttttttcaaaatttggtag
3775
225 I D N I D K A Y D L R K K I L N E F D K K C F L Q I W *

44AHJDORF009

5744 atgaaatcacacaacaagcaaaagaatggatatataagcatgagggggcaggtgttgactttgatgggtcatatggatttcaa
1 M K S Q Q Q A K E W I Y K H E G A G V D P D G A Y G P Q
5828 tgtatggacttatcagttgcttattgtgtattacattacggttaaaagttcgcattggtgggtaattgctaaagacgcgataaat
29 C M D L S V A Y V Y I T D G K V R M W G N A K D A I N
5912 aatgactttaaaggtttacgcagcgtgtatataaaatcacccgagctttaaaccctcaattaggggacgttgctgtatatataaat
57 N D F K G L A T V Y K N T P S F K P Q L G D V A V Y T N
5996 ggacaatatggacatatcaatgtgtgttaagtggaatcttgattattatatacatgcttagaacaacaaactgggttagcgcggt
85 G Q Y G H I Q C V L S G N L D Y Y T C L E Q N W L G G G
6080 tttagcggttgggaaaaagcaaccatttagaacacattattatgacggtgtaactcactttatttagacctaattttcaggtagt
113 F D G W E K A T I R T H Y Y D G V T H F I R P K F S G S
6164 aatagcaaaagcattagaacatcaaaagtaaatcatcttggaaaaatggaaacgaacaaacacgcgcacattattatagaatgaa
141 N S K A L E T S K V N T F G K W K R N Q Y G T Y Y R N E
6248 aatggtacattacatgtggtttttacaaatattgcacgtgtcggtagtccaaaattatcagaacctaaggtgattggttc
169 N G T F T C G F L P I F A R V G S P K L S E P N G Y F
6332 caaccaaacggttatcacattataaacgaagtttgtttatcagatggttacgtatggattggttataactggcaaggcacagct
197 Q P N G Y T P Y N E V C L S D G Y V W I G Y N W Q G T R
6416 tattattaccagtcgcgaatgggaatggaaaaacaggttaagttacagtggttggtattccttggtgggtggtctcctataa 6496
225 Y Y L P V R Q W N G K T G N S Y S V G I P W G V F S *

44AHJDORF010
14420 ttggttagacatacgtctgaaatggatagatggaaaaagaaagagaagctagaaaagagcaagaaaagattttatttttaaat
1 L V R H T S E M D R W K K E R E A R K E Q E K D L F L N
14336 gatttttagtaattgttaatttttaatttgatgataaagatttacaagaggctacattgacacatggaacattttgacacatcg
29 D F S N V N F K F D D K D L Q E A Y I D T W K H F A H L
14252 cctatttttctaagaagaagaaacgtatcatatgtaaatgctgtatcattggtaagagggttcaagacataaaaaatttaattat
57 P Y F P K E R N V S Y V N A V S L V R G S R H K K L N Y
14168 attcttgaataataataaccgtaattgatcttaataataaaaaacgctaaaaagcataaataacgctttatataatttacaagct
85 I L E I Y N R N D D S N K N A K K H K Y A L Y G A Q A
14084 aaaaataataatttcaatgtataaataatataaagaatcgatactttatataaagaatttgtaaatcagatagaccagtg
113 K N N N S S M Y K Y I K E I D T L Y K E I G K S D R P V
14000 acaaatattgatgaagatgtgaggataaactttttattattatgaacatttgacgaataa 13938
141 T N I D D E D V R Y N F L Y Y A T P D E *

44AHJDORF011
15593 atgacaaacgtaaaagatttttatcaagacaccaaacaacatttagcgagatttgaatttgaggaagaaagagaatttatc
1 M T N V K D I L S R H Q N T L A R F E F E E K E R E F I
15509 aaacttcagaatttagtagaaaaatacgggtatgaaaaagagtatatcggttagagcattattcacaaacaagaatcaaaattc
29 K L S E L V E K Y G M K K E Y I V R A L F T N K E S K F
15425 ggtgaacaagggtgttatcgctcactgatgactataacgtaaaacttaccgaaccacttaacagaatttaataagaagatgagca
57 G E Q G V I V T D D Y N V N L P N H L T E L I K E M R A
15341 gatgaggagcgtgttgacattatcaatgctggagaagttcaattcacattttatgaatatgaaaacaaaaagggtcaaaaagggt
85 D E D V V D I I N A G E V Q F T I Y E Y E N K K G Q K G
15257 tactcaatcaattttggtcaagtatcattttaa 15225
113 Y S I N F G Q V S F *

44AHJDORF012
8391 atgaacgaagtaaaattcagattttacagactcagaagcgtttcacatgtttatatacgcgtgggattttaaattactctacttt
1 M N E V K F R F T D S E A F F H M F I Y A G D L K L L Y F
8475 ttatttgaattaatgttcgttgatattattacagattttcaaaagcaattaaaaataaacttaattggtcgaagaaatcaatg
29 L F V L M F V D I I T G I S K A I K N N N L W S K K S M
8559 agaggattttctaaaaaattattgatattctgtattatcatttttagcaaacattcattgaccagattttacaattaaagggtggt
57 R G F S K K L L I F C I I I L A N I I D Q I L Q L K G G
8643 ctactcatgattacaattttttattatttgcataaggggactttctattgtagaaaattgtgcgaagtgacgtatttagta
85 L L M I T I F Y Y I A N E G L S I V E N C A E M D V L V
8727 ccagaacaaattaaagataaattagagtcattaaaaatgatactgaaaagagtgataacaatgaacgatcaagagaagataga
113 P E Q I K D K L R V I K N D T E K S D N N E R S R E D R
8811 taa 8813
141 *

44AHJDORF013
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1 M K I K T T F R L N N L I Y Y L L T N R D Y Y N D K F E
14912 aaatttacttcatctataaaaaatgtatagtaaaaaataatagggtgatgtgtatattgagtttgacaaacaatatgatg
29 K F T S S N K K C I V K I N M G D V Y I E F D K Q Y D D
14828 ttgaaattgaaaagagttattttacgttagatattcgacattgatattaaaaacatgtttttaaatactgtattttattat
57 F E I E K E L F T L D I D I D I K K H V F N I L V F Y Y
14744 agaaattatttaagtaattgaattaaagagaatttttattaaacgtttacaattgacgacgtattattcaaattttgataaacct
85 R N Y L S N E L I R E I L L N V T I D D V L S N F D K P
14660 cttgaaagcgaattaatgattttatcaaaacaaagtcataacgataatgggaaagtgattgacctgaataa 14586
113 L E S E L M I I Y Q N K V I Y D N G K V I D H E *

44AHJDORF113
199 atgacagaatttgatgaaatcgtaaaacccagcagcaagaagaacttcagaatcaactgaagaaaatttagaatcaactgaa
1 M T E F D E I V K P D D K E E T S E S T E E N L E S T E
283 gaaacttcagaatcaactgaagaatcaactgaagaatcaactgaagaatcaactgaagataaaacagtagaacaacacgaa
29 E T S E S T E E S T E E S T E E S T E D K T V E T I E E
367 gaaaatgaaaacaaattagaacctactacaacagatgaagatagttcgaaatttgacctgtgtattagaacaacgtattgct
57 E N E N K L E P T T T D E D S S K F D P V I A L
451 tcattagaacaacaagtgactactttttatcttcacaaatgcaacaaccacacaagtaacaacaacacaaatcagatgaaca
85 S L E Q Q V T T F L S S Q M Q Q P Q Q V Q Q T Q S D V T
535 gaatcaacaagaagataacgactattcagatgaagaactagttgataagttagatttagattag 600

277

113 E S N K E D N D Y S D E E L V D K L D L D *
44AHJDORF114
16172 atggttaattgttgataatgcaccagaagaaaaaggacaagcctatactgaaatgttgcaactattcaataaactgattcaatgg
1 M V N V D N A P E E K G Q A Y T E M L Q L F N K L I Q W
16088 aatccagcttatacatttgacaatgcaattaacttattatcggttgccaaactatttataaactataatagttctgtgtt
29 N P A Y T F D N A I N L L S A C Q Q L L L N Y N S S V V
16004 caattcttaattgatgaactaaacaacgaaactaaaccagaatcaatattgtcttatattgtctgtgatgaccaatagaacaa
57 Q F L N D E L N N E T K P E S I L S Y I A G D D P I E Q
15920 tggaatatgcataaaggattttatgaaacgtataacgtttacgttttttag 15870
85 W N M H K G F Y E T Y N V Y V F *
44AHJDORF014
6243 atgaaaatgggtacatttacatgtgtgtttttaccatattttgcacgtgtcggttagtccaaaattatcagaacctaatggctatt
1 M K M V H L H V V F Y Q Y L H V S V V Q N Y Q N L M A I
6327 gggtccaaccaaagggttatacaccatataacgaagttgtttatcagatgggttacgtatggattgggtataactggcaaggca
29 G S N Q T V I H H I T K F V Y Q M V T Y G L V I T G K A
6411 cacgttattattaccagtcgcgaatggaatggaacaggttaattgttacgtgtgtgtattccttggggggtgttctcat
57 H V I I Y Q C A N G M E K Q V I V T V L V F L G G C S H
6495 aatgggtatttttagcctttttctttga 6521
85 N G Y F S L F L *
44AHJDORF015
15403 gtgacgataacaccttgttcaccgaattttgattctttgtttgtgaataatgctctaactatactctttttcataccgtat
1 V T I T P C S P N F D S L F V N N A L T I Y S F F I P Y
15487 ttttctactaattctgatatgtttgataaattcttttttttctcctcaaatcaaatctcgctaatggtgtttgtgtctgtat
29 F S T N S D S L I N S L S F S S N S N L A N V F W C L D
15571 aaaatatctttttagcgtttgtcatttttttctcctctattttaaattattgttttctgcaattgcgattgttag 15645
57 K I S F T F V I L F L L L F K L F A F C N C D L *
44AHJDORF016
15852 atgaaagttgacgacattgttaccttacctgtcgaaggttatatacttctacttagatgatgataatgaatacattgaggaa
1 M K V D D I V T L R V K G Y I L H Y L D D D N E Y I E E
15768 ttttaccacttcacgagtatcattttaacaaaacacaaagcaaaatattaccagacacatgtaaaactattgtccactaca
29 F L P L H E Y H L T K T Q A K E L L P D T C K L L S T T
15684 cgcacaacgaaaacattcaagttattacaatgatttactacaaatcgcaattgcagaagcaataa 15616
57 R T T K T I Q V Y Y N D L L Q I A I A E S K *
44AHJDORF017
10757 atgaaagattaaaattgcttctgctgtgtataccgaaaacgccttttgatacaagcgtcgattttgaaacctttgtacgtgaac
1 M E L K L L L V Y R K T P L I Q A S I L K P L Y V N
10673 aattctttgacgggtgccattattgaaaacaataaaagtattctataatgagcaagggtacaatatcgatatatccgtctaaaactg
29 N S L T V P L L K T I K V S I M S K V Q Y R Y I R L K L
10589 aaattgtatgtgtgaatgtatatgatgaatattttactgatgaacttaatatga 10536
57 K L Y V V M Y M M N I L L M N L I *
44AHJDORF018
1098 atgttaattggtactgtgtccataatcacgtattcttctcactatattgtccaataaaatcttgctcttttagctaaccaattaaaa
1 M L I G T V S I I T Y S S L Y C P I K S C S L A N Q L K
1014 cgattacctaattgcaatatcgattaataaagtctcattatcttagggaataaatattttattacaaatgtttcgaaacttgta
29 R L P N A I S I N K V S L I L G N K Y L F T N V S N I V
930 tttgattatcccatgtgtcgcaaatgtccaagattttgaataa 886
57 P E L S H L S P N V Q D F E *
44AHJDORF019
9836 atgttacctggtttgtataagattcttttttgaataaagggtacaccaattgcttttttatatttttctggttaactgtgcatat
1 M L P G L Y K Y S F L N K G T P I A F L Y F S G N C A Y
9752 gtccagttaccacaaatcacagaccactttttccattgtggttgactgattaccactaattgggttatgggtctccgtcatca
29 V Q L P P I T R P L P P F G L T D L P L I G L W S P S S
9668 tcagtaggattagaactactactccactatctactga 9630
57 S V G L E L L L P L S T *
44AHJDORF121
16362 atggaaaatgaacaaaaaacattgagttgaagcatgttttctgttttaagaatggaagtttatgtatagcgttatttgataga
1 M E N E T K N I E L K H V F R F K N G S L C I A L F D R
16278 acagaaaatgaatttctatttatgtatgttgacattgatgaattgaagatttaaatcataattctgttttacgcgtaatttca
29 T E N E I S F Y D V D I D E I E D L N H N S V L R V I S
16194 actttatttaggaagtataaataatggttaa 16165
57 T L L G S D N N G *
44AHJDORF020
13865 atgtctaaacgattttgttttaccatgtttttgtcctctgtaaatagtttatgatgtcggtttacagtgttaattttattcgctcaa
1 M S K R F C F T M F L L L V I V Y D V V Y S V K F I R Q
13949 atgttgataataaaaaagttatacctcacatcttcatcatcaatattgtcactggtcttatctgatttaccatttctttat
29 M L H N I K S Y T S H L H H Q Y L S L V Y L I Y Q F L Y
14033 ataaagtatcgattttcttaa 14053
57 I K Y R F L *
44AHJDORF123
614 atgtatgagggaaacaacatgcgttcttatgatgggtacatcatatgaagattcaagattaaataaacgaacagaattaaatgaa
1 M Y E G N N M R S M M G T S Y E D S R L N K R T E L
698 aacatgtcaattgatacaataaaagtgaagattgtgtgtacaaattcattcactttcaaaacaatcatttacaggtgac
29 N M S I D T N K S E D S Y G V Q I H S L S K Q S F T G D
782 gttgagggaggaataa 796
57 V E E E *

44AHJDORF021

5816 atgcaccatcaaagtcaacacctgccccctcatgcttatatatccattcttttgcttgtgtgtgatttcatttatatcactc
1 M H H Q S Q H L P P H A Y I S I L L L V V V I S P I S L
5732 ctatttttgatgttttgctacccaaccatattcacgatgttttgcttccgcattacattactgaagaattctttatattccga
29 L F L M F C Y P T I F T M F C F R I N I T E E F F I F R
5648 tatattagcctctaa 5634
57 Y I S L *

44AHJDORF022

8611 atgtttgctaaaatgataatacagaatatcaataatttttagaaaatcctctcattgattttttgaccataagttattattt
1 M F A K M I I Q N I N N F L E N P L I D F F D H K L L F
8527 ttaattgcttttgaaatccctgtaataatatcaacgaacattaatacaaaaaaagtag 8468
29 L I A F E I P V I I S T N I N T N K K *

44AHJDORF023

6494 atgagaacaccccccaaggaataccaacactgtaactattacctgtttttccattccattggcgactggtaataataacgtg
1 M R T T P P K E Y Q H C N Y Y L F F H S I G A L V N N N V
6410 tgctttgcccagttataaccaatccatcacgtataaccaacttcggttatatgggtgtataaccggtttgggtggaacc
29 C L A S Y N Q S I R N H L I N K L R Y M V Y N R L V G T
6326 aatagccattag 6315
57 N S H *

44AHJDORF024

14275 gtgtcaatgtacgcctcttgtaaatctttatcatcaaatttaaaattaacattactaaaatcatttaaaaaataaatctttttct
1 V S M Y A S C K S L S N L K L T L L K S F K N K S F S
14359 tgctctttttctagcttctctttctttttccatctatccatttcagacgtatgtetaaccaatgttatcaacctccatataaag
29 C S F L A S L S F F H L S I S D V C L T N V I N L H I K
14443 cataataa 14451
57 H K *

44AHJDORF025

15175 atggaacgtaaatacaaaaacggtattatttatattgcgatgagattaaggacattttccacatcaaatctcaatgtttgaagat
1 M E R K Y K T V L L Y C D E I K G H F P H Q I S M F E D
15091 ttatatgacgctaaagtgtatattcatattatgaatataacctgttcactaaaaatacgcgtatatacatagaatacattaag
29 L Y D A K V V Y S Y Y E Y N L F T K K Y A Y I I E Y I K
15007 gagatataa 14999
57 E I *

44AHJDORF026

14593 atgaataacctattaaacatagccattgttttcttttagcatttttaattacattatcatacttatgacactgcatatacgc
1 M N N L L N I A I V F L L A F L I T L I I L M T L H I R
14509 gtgtcatttggtgttttattcactacattgattatattctatattatcttttaagtgttatttatgctttataggagggtga
14426
29 V S F G V L F T T L I I F Y I I F L M V I Y A L Y G G *

44AHJDORF027

12916 atgattgtctatatccctaatttttagtataaaattcatattgttttgatatgggtacaacgataatattttgcataaaagtagt
1 M I V Y I P N F S T K F I L F C I W Y N D N I C H K S S
13000 tacattatacatgactttaatatattatcatcagttttgatataagaagaatcaccgttttgattgatgtgatttcttaa
13080
29 Y I I H D F N I F I I S F D I E E I T V L I D V I S *

44AHJDORF029

15183 gtgttttaaatggaacgtaaatacaaaaacggtattatttatattgcgatgagattaaggacattttccacatcaaatctcaatgt
1 V F K W N V N T K R Y Y Y I A M R L K D I F H I K S Q C
15099 ttgaagatttatatgacgctaaagtgttatattcatattatgaatataacctgttcactaaaaatacgcgtatatacatg
15019
29 L K I Y M T L K L Y I H I M N I T C S L K N T R I S *

44AHJDORF028

9235 atggaatatatgcacgtccaattgtacctgctttcatattttttgcaaaatctgcattaccttttctttgtacgtcttggta
1 M E Y M H V Q L Y L L S Y F L Q N L H Y L F F V R L V V
9151 caaagtggacgatgttacctgcgtcatccaagacggtgtgccagcttgttttgattgtgataacttcttctgctatga 9071
29 Q S G R C Y L R H T K T V V Q L V L I V I L T F L L *

44AHJDORF030

14487 gtgaataaaacaccaaataacacgcgttatatgcagtgctataagtgataagtgtaattaaaaatgctaaaaggaaacaatg
1 V N K T T P N D T R I C S V I S M I S V I K N A K R K T M
14571 gctatgtttaatagggttattcatggtcaatcactttccattatcgatatgactttgttttgataaaataatcattaa 14648
29 A M F N R L F M V N H F P I I V Y D F V L I N N H *

44AHJDORF031

11039 atgatattgtatagttcattgttatctctaaacggaataagtgtaaaatgtgaacgtaaatgcaggtatgccatataatccattt
1 M I L Y S S L L S S K R N K L K C E R N A G M P Y N P F
11123 aaaacgacttttagataacataacctcctcattttgagtgtgtgttcgttgatatcatcagtaaatgtga 11191
29 K T T L D N I T S S F E Y G C S L I S S V M *

44AHJDORF035

693 atgaaaacatgtcaattgatcaaaaatgaagtgaagatagttatgggtgacaaattcattcactttcaaaacaatcattacag
1 M K T C Q L I Q I K V K I V M V Y K F I H F Q N N H L Q
777 gtgacgttgaggaggaataataaattatggcacacaatactacaaaaatgaaactgcacttttag 842
29 V T L R R N N K L W H N N L Q K M K L H F *

44AHJDORF033

3795 atgccattatttaaccacctctaccaaatttgtaaaaaacattttttatcaaattcatttaaaattttctttcttaaatcgta
1 M P L F N H L Y Q I C K K H F L S N S F K I F F L K S Y

3711 gctttatcaatattatcaattaaactgcttagtggaattgtgtaccttttgcattaccttttga 3646
29 A L S I L S I K Y C L V N C V P F A L P F *
44AHJDORF032
9455 atggctgtgttttctaaagcgagtagtgaaactaccactgtcaccactactaccactgtcagacgaatcactaggtgatccacct
1 M A C F A K A S S E L P L S P L L P L S D E S L G D P P
9371 ttaccgtctaatttaccacccaagctagaatagatttcgaccgtctaaaaatggattaccatag 9306
29 L P S N L P P Q A R I V F A P S K N G L P *
44AHJDORF034
14146 atgatgattcttaataaaaaacgctaaaaagcataaatacgttttatataatttacaagctaaaaataataattcttcaatgt
1 M M I L I I K T L K S I N T L Y I I Y K L K I I I L Q C
14062 ataaatatattaagaaatcgatactttatataaagaaattggtaaatcagatagaccagtga 14000
29 I N I L K K S I L Y I K K L V N Q I D Q *
44AHJDORF035
13957 atgcaacatttgacgaataaatttaacactgtaaacgacatcataaactattacaaggagcaaaaacatggtaaaacaaaatcg
1 M Q H L T N K F N T V N D I I N Y Y K E Q K H G K T K S
13873 tttagacatggtgaagattatcaaatgctgtcaatcatgtcagaaaaaaatcccagataa 13811
29 F R H G K R L S K C C Q S C Q K K N P R *
44AHJDORF036
10165 gtgtatacaataccacacgtgatggtgcaacatattggtgtacattatagtttgcaactaaaaacgaaccatcttcaaaaactg
1 V Y T I P H V M V Q H M V V H Y S L Q L K T N H L Q K L
10081 ctacaacaacacctgtgtgaccaataccatattgcagttgcttgaagtatggtggtttactag 10019
29 L Q Q H L C D Q Y H M Q L L V S M V V Y *
44AHJDORF037
14788 atgtcgatatctaacgtaataaactctttttcaatttcaaaatcatcatattggttgcactcaatatacacatcacccata
1 M S I S N V N N S F S I S K S S Y C L S N S I Y T S P I
14872 tttatttttactatacattttttattagatgaagtaatttttcaattttatcattataa 14931
29 F I F T I H F L L D E V N F S N L S L *
44AHJDORF038
3671 gtgtaccttttgcattacctttttgattttgattacgtttttgcttttgcattactttcgttactcgattttacagttttac
1 V Y L L H Y L F D F D Y V L R F D Y F R Y S I Y S Q F Y
3587 cgttatcaatcgattatttatcagcgaatcgtaacgttgtattatcaacatcaatgttaa 3528
29 R Y Q S Y Y Y Q R I V T L Y Y Q H Q C *
44AHJDORF039
1743 gtgctgtatttacttatgatgtatctaaacttaagagtttactggcaacgttgaagaaattaaacccaaatcagatttatag
1 V L Y L L M M Y L N L K S L L A T L K K L N Q N Q I Y M
1827 cgtttattttggatattaattcaattataaaacgttacacaaaagggtatgttaa 1883
29 R L F W I L I Q L N I N V T Q K V C *
44AHJDORF040
9740 gtggaactggacatattgcacagttaccagaaaaatataaaaaagcaattggtgtacctttattcaaaaaagaatacttatata
1 V V T G H M H S Y Q K N I K K Q L V Y L Y S K K N T Y T
9824 aaccaggtaacatatttctcctcaaacgggtaatgcaggacaatgtacagaattaa 9877
29 N Q V T Y F L K R V M Q D N V Q N *
44AHJDORF041
15836 atgtcgtcaacttttcattattatatactccttttctaaaaacgtaaacgtttatcagtttcataaaatcctttatgcattatcc
1 M S S T F I I I S L L S K K R K R Y T F H K I L Y A Y S
15920 attgttctatttgggtcatcaccagcaatataagacaatattgattctcgttttag 15973
29 I V L L G H H Q Q Y K T I L I L V *
44AHJDORF042
5151 atgcacgaccgtcgtctttttgttaatttatagttttgtgaacctcttgcgcgtaatgcttcaaaagtgttcataactaccaagtt
1 M H D R R L L L I Y S F V N L L R V M L Q S V H T H Q V
5067 ggaagaaaccataataaattatggaacggttttccaccacgcggtttgtcatag 5014
29 G R N H I N Y G N V F H H R R L S *
44AHJDORF043
4539 atgcgacttgtaacagttttgcaacacatcgtagtaaccagattttcatttcaccattggattgacgttctaatccgattg
1 M R L V T V L Q H H R D V T R F S F H H W I D V L I R L
4455 ttgtaccatgaccaccctgtacaatacgcagcttgaaattaaagtcaccactag 4402
29 L Y H D H P V Q Y A C L K L S H H *
44AHJDORF044
12917 atgttacctattttacgtgatgatattttataaagaaaaacatggaacgttattactacaatccaagcaattttacattttgaca
1 M L P I Y V M I C F I K K T W N V I T T I Q A I Y I L T
12833 atgcttactctaaaaattacgtggttgataatgatagatatttatatttag 12783
29 M L T L K I T W L I M I D I Y I *
44AHJDORF149
770 atgattgtttgaaagtgaatgaattgttacaccataactatcttcactttttattgtatcaattgacatgttttcatttaatt
1 M I V L K V N E F V H H N Y L H F Y L Y Q L T C F H L I
686 ctggttcgtttatttaattcttgaatcttcattatgatgtacccatcatag 639
29 L F V Y L I L N L H M M Y P S *
44AHJDORF046
4891 atgattatccatttaagtattcatatcaagacggtatttaatttccacgtgataaacttttaagagcctgaagggtatttgcattt
1 M I I H L S Y H I K T V L I S H V I T L K S L R V F A F
4975 atacaaatccaaaaacaaacgtaaatcgttattacttgcattatga 5019
29 I Q I Q K Q N V N R Y Y L L *
44AHJDORF047
11911 atgaatgtatgaagtgttcaggtgtgagttttgcaaacattttcacagcatagtcaggttctactatcattcatattcatt
1 M N V C K L F R C E F C K T F H S I V I G F T I I H I I

11995 atctttatcaaaaatcgatataattaaaatctgttttaagtgtga 12039
29 I F I K N R I I K I C F K L *
44AHJDORF045
10655 atggcaccgtcaaagaattgttcacgtacaaagggtttcaaaatcgacgttgatcaaaggcgttttcggtataaccagcagaa
1 M A P S K N C S R T K V S K S T L V S K A F F G I P A E
10739 gcaattttaatctttccattcacttcattatgcataattcttatga 10783
29 A I L I F P P T S Y A Y F L *
44AHJDORF048
15340 atgaggacgtgttgacattatcaatgctggagaagttcaattcacaatttatgaatatgaaaacaaaaagggtcaaaaagggt
1 M R T L L T L S M L E K F N S Q P M N M K T K K V K K V
15256 actcaatcaatttttggtcaagtatcattttaatacaatttcatag 15212
29 T Q S I L V K Y H F N T I S *
44AHJDORF049
5784 atgagggggcaggtgttgactttgatgggtgcataatgatttcaatgtatggacttatcagttgcttatgtgtattacattactg
1 M R G Q V L T L M V H M D F N V W T Y Q L L M C I T L L
5868 acggtaaagttcgcattgtgggtaaatgctaaagcgcgataa 5909
29 T V K F A C G V M L K T R *
44AHJDORF050
13158 gtgtgttactgttttcattcacgtaatcgtttcgtcgatttctaaaaaatgtttttgtaaagtcttgatgtattcattttat
1 V C Y V F H S R N R F V A F L K K C F C K V L M Y S F Y
13242 gctttttgtaataaattgtatatatttaattggataatag 13283
29 A F V I N C I Y L N W I I *
44AHJDORF051
11066 atgataacaatgaactatacaatcattaacggttacaaaaactgaacgtaatatattattctctacatttgcacatcac
1 M I T M N Y T I S L T V T K T L N V I Y Y S L H L S H H
10982 gttcattgtataacttattggttcccttccaatacttaa 10944
29 V H C I T Y W F L S N T *
44AHJDORF052
14338 atgatttttagtaattgttaatttttaattttgatgataagatttacaagaggcgtacattgacacatggaaacattttgcacatc
1 M I L V M L I L N L M I K I Y K R R T L T H G N I L H I
14254 tgccctattttcctaagaagaacgtaacatgataa 14216
29 C P I F L K K E T Y H M *
44AHJDORF053
3348 atgtgggtttattcatcaagtgaagttgaaaaatacttacaatcacaaggcgttcacagaacacaaatgaagatacaacaagtaaca
1 M W F I H Q V K L K N T Y N H K A S Q N T M K I Q Q V T
3432 ctgatgaacatcgaatcaaatgctacatctttag 3467
29 L M K H R I K M L H L *
44AHJDORF054
7551 atgactggaatggaaatcagatgttactcgacgtcgtgaagatttcacaaaaactgggtgtaagttacgtacaaaatcaatta
1 M T G M E I R C Y S T L V R F H K K L V L S Y V Q N Q L
7635 ttggttatcataatgaagttcgagtatatccagtag 7670
29 L V I I M K F E Y I Q *
44AHJDORF055
15705 atgtgtctggtaataattcttttctgtgttttggttaaatgataactcgtgaagtggtaaaaattcctcaatgtattcattat
1 M C L V I I L L L V F W L N D T R E V V K I P Q C I H Y
15789 catcatctaagtaagtaagtatataacctttga 15821
29 H H L S N E V Y N L *
44AHJDORF056
5512 gtgagttattacattacaggttaacaaatggaattatttagagacgcgcgacagaagaataaaaaagggtgggtgcattggtacgtg
1 V S I T L Q V T K W N Y L E T R Q K K L K K W V H G Y V
5596 tgtcaagtggttaacgcagtcggtgaagtaa 5625
29 C Q V V T Q S V K *
44AHJDORF057
10121 atgtaccaccatattgttcaccatcagtggtgtattgtataactcattaatggcgtaccaaataatgctgggtgataatattg
1 M Y H H M L H H H V W Y C I H S L M A Y Q I M L V I I L
10205 tattcttttagtggtattgcttaattaa 10231
29 Y S L V V L L N *
44AHJDORF058
10767 atgcataattttcttatgattcagtaacaacatcttatctctgttctgttttcaatatccatttacctaaggctatcggtcga
1 M H I S Y D S V Q T S Y L S V R F Q Y P I Y L R L S G R
10851 ataaactgggttcaataagggtttaa 10877
29 I N W G S I R V *
44AHJDORF164
702 atgttttcatttaattctgttcgtttattttaattcttgaattcttcatatgatgtacccatcatagaacgcattgtgtttccctca
1 M F S F N S V R L F N L E S S Y D V P I I E R M L F P S
618 tacatgttttaattcctcctaatactaa 592
29 Y M F K F L L I *
44AHJDORF059
8360 atggattttgtaacattggattacgtgaacggtcattatgccaaatcttacaccagattctaaaattgcttttaattgttcca
1 M D F V T L D Y L N R H Y A K I L H Q I L K L L L I V P
8276 ttaacatgggtcgtatgtcacgtatag 8250
29 L T W G R C H V *
44AHJDORF060
6257 atgtaccattttcattttctataatattgtccgtattggtttcgtttccattttccaaatgtatttacttttgatgttttctaag
1 M Y H F H F Y N M C R I G F V S I F Q M Y L L L M F L M

281

6173 ctttgctattactacctgaaaatttag 6147
29 L C Y Y Y L K I *
44AHJDORF061
15551 atgtgttttgggtgtcttgataaaaatcttttacggttgctattttatttctcctcttatttaaattatttgctttctgcaatt
1 M C F G V L I K Y L L R L S F Y F S S Y L N Y L L S A I
15635 gcgattttagttaaattcattgtaa 15658
29 A I C S K S L *
44AHJDORF062
14285 gtggtattcgcaacgcagttaaccaatctattaatattgataaagaacaaatcacatgtactctacacaatccgattctcaaa
1 V V F A T Q L T N L L I L I K K Q I T C T L H N P I L K
14369 aacctgaagggttttggataa 4389
29 N L K V F G *
44AHJDORF063
19487 atgcgtcttgatttttttaataattcttgcattggcttgttttgctaaagcgagtagtgaactaccactgtcaccactactac
1 M R L V F F L I I L A W L V L L K R V V N Y H C H H Y Y
19403 cactgtcagacgaatcactag 9383
29 H C Q T N H *
44AHJDORF065
15029 gtggtggaaaaacgtttccataatttatatgggtttcttccaacttgggtgagtatgaacactttgaagcattacgcgcaagagggt
1 V V E N V S I I Y M V S S N L V S M N T L K H Y A Q E V
15113 cacaaaactataaattaa 5130
29 H K T I N *
44AHJDORF064
12609 atgacgagtcaatcaatcaacttgtgtccgaaatatataacggtgcaccatttgttaaaatgtcacctatgtttaatgcagatg
1 M T S Q S I N L C P K Y I T V H H L L K C H L C L M Q M
12693 acgatatcattgatttaa 2710
29 T I S L I *
44AHJDORF066
110481 atgatattctttatattgaaagtgcacatcggttcattttcacttaacgacttatttccagttgaacgttcagtacataacaaat
1 M I F F I L K V T S V H F H L T T Y F Q L N V Q Y I T N
110397 ctgatttgcataatattaa 10380
29 L I C I Y *

Table 19

Sequence similarities between ORFs 44AHJD and public databases

Phage: 44AHJD

Database: nr

Query= sid|110871|lan|44AHJDORF001 Phage 44AHJD ORF|10342-12627|-1
(761 letters)

gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE >gi 76896 pir JQ0...	55	1e-06
gi 1072656 pir S51275 DNA polymerase - phage CP-1 >gi 836593 e...	53	6e-06
gi 1429230 emb CAA67649 (X99260) DNA polymerase [Bacteriophage...	49	1e-04
gi 1572479 emb CAA65712 (X96987) DNA polymerase [Bacteriophage...	46	0.001
gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...	45	0.002
gi 2435429 (AF012250) unassigned reading frame (possible DNA po...	45	0.002
gi 1084487 pir S41618 DNA polymerase - slime mold (Physarum po...	45	0.002
gi 4877819 gb AAD31446.1 (AF133505) DNA polymerase [Neurospora...	44	0.004
gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE >gi 2833...	44	0.004
gi 2499511 sp Q12471 6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2 (PHO...	41	0.041
gi 2258375 gb AAD11909.1 (AF007261) transcription initiation f...	40	0.070
gi 15734 emb CAA37450 (X53370) DNA polymerase (AA 1-575) [Bact...	39	0.092

Query= sid|110872|lan|44AHJDORF002 Phage 44AHJD ORF|3789-5732|3
(647 letters)

gi 135273 sp P27622 TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTE...	112	7e-24
gi 142847 (M64050) DNase inhibitor [Bacillus subtilis]	52	1e-05
gi 4038407 (AF103943) factor C protein precursor [Streptomyces ...	39	0.10

Query= sid|110873|lan|44AHJDORF003 Phage 44AHJD ORF|6626-8389|2
(587 letters)

gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...	92	8e-18
gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...	82	1e-14
gi 1429238 emb CAA67657 (X99260) tail protein [Bacteriophage B...	78	2e-13
gi 215339 (M12456) p9 tail protein [Bacteriophage phi-29] >gi 2...	71	2e-11
gi 1181968 emb CAA87738.1 (Z47794) tail protein [Bacteriophage...	54	3e-06
gi 1181970 emb CAA87740.1 (Z47794) tail protein [Bacteriophage...	42	0.010

Query= sid|110875|lan|44AHJDORF005 Phage 44AHJD ORF|12643-13890|-1
(415 letters)

gi 3845203 (AE001399) GAP domain protein (cyclic nt signal tran...	52	6e-06
gi 3758843 emb CAB11128.1 (Z98551) predicted using hexExon; MA...	49	5e-05
gi 3845297 (AE001421) hypothetical protein [Plasmodium falciparum]	48	1e-04
gi 4493936 emb CAB38972.1 (AL034556) predicted using hexExon; ...	47	2e-04
gi 3845165 (AE001390) hypothetical protein [Plasmodium falciparum]	46	6e-04

Query= sid|110877|lan|44AHJDORF007 Phage 44AHJD ORF|2044-3027|1
(327 letters)

gi 1181960 emb CAA87731.1 (Z47794) connector protein [Bacterio...	46	5e-04
gi 1429239 emb CAA67658 (X99260) upper collar protein [Bacteri...	45	8e-04
gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...	44	0.002
gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...	41	0.009

Query= sid|110878|lan|44AHJDORF008 Phage 44AHJD ORF|3020-3775|2
(251 letters)

gi 4982468 gb AAD30963.2 (AF118151) SNF1/AMP-activated kinase ...	52	3e-06
gi 1730077 sp P18160 KYK1_DICDI NON-RECEPTOR TYROSINE KINASE SP...	46	2e-04
gi 3758855 emb CAB11140.1 (Z98551) predicted using hexExon; MA...	46	2e-04
gi 585795 sp P21538 REB1_YEAST DNA-BINDING PROTEIN REB1 (QBP) >...	46	3e-04
gi 172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]	46	3e-04
gi 2952545 (AF051898) coronin binding protein [Dictyostelium di...	45	6e-04
gi 535260 emb CAA82996 (Z30339) STARP antigen [Plasmodium reic...	45	7e-04
gi 1429240 emb CAA67659 (X99260) lower collar protein [Bacteri...	44	0.001

Query= sid|110879|lan|44AHJDORF009 Phage 44AHJD ORF|5744-6496|2
(250 letters)

gi 2764981 emb CAA69021.1 (Y07739) N-acetylmuramoyl-L-alanine ...	180	1e-44
gi 113675 sp P24556 ALYS_STAAU AUTOLYSIN (N-ACETYLMURAMOYL-L-AL...	118	6e-26
gi 1763243 (U72397) amidase (bacteriophage 80 alpha)	118	6e-26
gi 4574237 gb AAD23962.1 AF106851.1 (AF106851) LytN [Staphyloco...	84	9e-16
gi 3767593 dbj BAA33856.1 (AB015195) LytN [Staphylococcus aureus]	84	9e-16
gi 2764983 emb CAA69022.1 (Y07740) cell wall hydrolase Ply187 ...	77	2e-13
gi 3287732 sp O05156 ALE1_STACP GLYCYL-GLYCINE ENDOPEPTIDASE AL...	73	2e-12
gi 79926 pir A25881 lysostaphin precursor - Staphylococcus sim...	69	3e-11
gi 126496 sp P10548 LSTP_STAST LYSOSTAPHIN PRECURSOR (GLYCYL-GL...	69	3e-11
gi 3287967 sp P10547 LSTP_STASI LYSOSTAPHIN PRECURSOR (GLYCYL-G...	69	3e-11
gi 3341932 dbj BAA31898.1 (AB009866) amidase (peptidoglycan hy...	68	6e-11

Query= sid|110882|lan|44AHJDORF012 Phage 44AHJD ORF|8391-8813|3
(140 letters)

gi 140528 sp P24811 YQXH_BACSU HYPOTHETICAL 15.7 KD PROTEIN IN ...	80	6e-15
gi 4126631 dbj BAA36651.1 (AB016282) ORF45 (bacteriophage phi-...	76	1e-13
gi 141088 sp P26835 YNGD_CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN ...	61	4e-09
gi 2293160 (AF008220) YtkC [Bacillus subtilis] >gi 2635548 emb ...	36	0.099
gi 1181973 emb CAA87743.1 (Z47794) holin protein [Bacteriophag...	31	3.3

Table 20

Homologies between phage 44 AHJD ORFs and proteins in public databases

Query= pt|110871 44AHJDORF001 Phage 44AHJD ORF |10342-12627|-1 1
(761 letters)

>gi|118848|sp|P19894|DPOL_BPM2 DNA POLYMERASE >gi|76896|pir||JQ0161
DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2
>gi|215509 (M33144) DNA polymerase (Bacteriophage M2)
Length = 572

Score = 55.4 bits (131), Expect = 1e-06

Identities = 96/426 (22%), Positives = 159/426 (36%), Gaps = 88/426 (20%)

Query: 229 KLTPEQLTYIHNDVIIIGMCHIHYSDFPNFDYNKLTFSNLNIMESYLNEMTR-----FQ 283
++TPE+ YI ND+ I+ DI +++T + + + + T+ F
Sbjct: 154 EITPEEYIKNDIEIARA----LDIQFKQGLDRMTAGSDSLKGFKDILSTKFNKVFP 209

Query: 284 LLNQYQDIKISYTHYHFDNMFDYDIKSFYRGGLNMYNTKYINKLIDEPFCSIDINSSYP 343
L+ D +I + YRGG N KY K I E D+NS YP
Sbjct: 210 KLSLPMDEI-----RKAYRGGFTWLNDRKYKEIGEGMV-FDVNSLYP 252

Query: 344 YVYHEKIPTWLYFYEHYSEPTLIPTFLDDNYFSLYKIDKDVFNDDLLIKIKSRVLRQM 403
MY +P Y P + + D + LY I + F +L K + +
Sbjct: 253 SQMYSRPLP-----YGAPIVFQGGYKDEQYPLY-IQRIREFEFEL---KEGYIPTI 299

Query: 404 XXXXXXXXXXXXXXXXXXXXLRMIQ-DITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
+ ++ +T +D I+ + + +Y EY F +
Sbjct: 300 QIKKNPFKNGEYLNKSGVEPVELYLTNVDLELIQEH-YELYNVEYIDGFK-----FRE 352

Query: 463 TQGLKLNKNIMTSPDYHITDDINEHPYSNEEVMLSQVVLNGLYG-----IPAL 511
G K+ I+ + H + L+K++LN LYG +P L
Sbjct: 353 TGLFLFDKDFIDKWTYVKT-----EAGAKQLAKLMLNSLYGKFASNPDPVTGKVPYL 403

Query: 512 RSHFNL-PRLDNNELNYIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDDNF 570
+ +L FR+ D YK+ + F+T+ + + + Q D
Sbjct: 404 KDDGSIGFRVGDDE-----YKDPVYTPM-GVFITAWARFTTITAAQACY-----DRI 449

Query: 571 IYCDTDSLVMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHK-----YAYEVNG 625
IYCDTDS+++ P + + DP LG W E+ + L K Y EV+G
Sbjct: 450 IYCDTDSIHLTGTEVPEIIKDIVDPKKLGYWAHES-TFKRAKYLRQKTYIQDIYVKEVDG 508

Query: 626 KIKIAS 631
K+K S
Sbjct: 509 KLKECS 514

>gi|1072656|pir||S51275 DNA polymerase - phage CP-1
>gi|836593|emb|CAA87725.1| (Z47794) DNA polymerase
(Bacteriophage CP-1)
Length = 568

Score = 53.5 bits (126), Expect = 6e-06

Identities = 104/464 (22%), Positives = 169/464 (36%), Gaps = 66/464 (14%)

Query: 230 LTPEQLTYIHNDVIIL--GMCHIHYSDFPNFDYNKLTFSNLNIMESYLNEMTRFQLLNQ 287
+ PE + YIH DV IL G+ ++Y + F Y + +L + +F+
Sbjct: 152 IKPEWIDYIHVDVAILARGIFAMYEEENFTK--YTSASEALTEFKRIFRKSRRKFRDFFP 209

Query: 288 YQDIKISYTHYHFDNMFDYDIKSFYRGGLNMYNTKYINKLIDEPFCSIDINSSYPVYMY 347
D K+ D+ + G + K+ + + + DINS YP M
Sbjct: 210 ILDEKVD-----DFCRKHIVGAGRLPTLKHRGRTLNQLIDIYDINSMPATML 257

Query: 348 HEKIPTWLYFYEHYSEPTLIPTFLDDNYFSLY-KIDKDVFNDDL-LIKIKSRVLRQMXX 405
+P + + Y P + +D+Y+ + K D D+ L I+IK ++
Sbjct: 258 QNALPIGIP--KRYKGG--PKEIKEDHYIYHIKADFLLKRGYLPITIKKKLDALRIG 312

Query: 406 XXXXXXXXXXXXXXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTG 465
L + + H + E F +F +Y
Sbjct: 313 VRTSDYVTTSKNEVIDLYLTNFDLFLKHYDATIMYVETLE-FQTSDFDDYI----- 366

Query: 466 KLKKNKINMTSPYDYHITDDINEHPYSNEEVMLSQVVLNGLYGIPALR--SHFNLFRLLDN 523
 + Y Y E+ S E +K++LN LYG + S L LDD
 Sbjct: 367 -----TTYRYK-----KENAQSPAOKQAKIMLSLYGKFGAKIISVKKLAYLDDK 412

Query: 524 NELYNIINGYKNTERNIL-----FSTFVTSRSLYNLLVPPQYLTESEIDNFIYCDTDS 577
 L +KN + + + FVTS + + + Q E DNF+Y DTDS
 Sbjct: 413 GILR-----FKNDDEEVQPVYAPVALFVTSIARHFIISNAQ-----ENYDNFLYADTDS 462

Query: 578 LYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHKKYAYEVNGKIKIASAGIPKN 637
 L++ +L+ DP GKW E + K L K Y E+ + + K
 Sbjct: 463 LHLFHSDSLVDL---IDPSEFGKWAHEGRAV-KAKYLRSLKYIEELIQEDGTTTHLDV-KG 517

Query: 638 AFDTSVDFETFFVREQFFDGAIIENNKSIYNEQGTISIYPSKTEI 681
 A T E E F GA E ++ +G IY + +I
 Sbjct: 518 AGMTPEIKEKITFENFVIGATFEGKRASKQIKGGTLIYETTFKI 561

>gi|1429230|emb|CAA67649| (X99260) DNA polymerase [Bacteriophage
 B103]
 Length = 572

Score = 49.2 bits (115), Expect = 1e-04
 Identities = 93/422 (22%), Positives = 155/422 (36%), Gaps = 88/422 (20%)

Query: 229 KLTPEQLTYIHNDVILGMCHIHYSDFPNFDYNKLTFSLNIMESYLNEMTR-----FQ 283
 ++TPE+ YI ND+ I+ DI +++T + + + + T+ F
 Sbjct: 154 EITPEEYIKNDIEIARA-----LDIQFKQGLDRMTAGSDSLGFKDILSTKKFNKVP 209

Query: 284 LLNQYQDIKISYTHYHFDNMNFYDIKSFYRGGLNMYNTKYINKLIDEPFCSIDINSSYP 343
 L+ D +I + YRGG N KY K I E D+NS YP
 Sbjct: 210 KLSLPMDEI-----RRAYRGGFTWLNDKYKEKEIGGMV-FDVNSLYP 252

Query: 344 YVMYHEKIPTWLYFYEHYSEPTLIPTFLDDNYFSLYKIDKDVFNDDLLIKISRVLRQM 403
 MY +P Y P + + D + LY I + F +L K + +
 Sbjct: 253 SQMYSRPLP-----YGAPIVFQGYEKDEQYPLY-IQIRFEFEL-----KEGYIPTI 299

Query: 404 XXXXXXXXXXXXXXXXXXXXLRMIQ-DITGIDCMHIRVNSFVIYCEYFHARDIIFQNYFIK 462
 ++ +T +D I+ + + +Y EY F +
 Sbjct: 300 QIKKNPFFKGNEYLKNSGAEPVELYLTNVDLELIQEH-YEMYNVEYIDGPK-----FRE 352

Query: 463 TQGLKKNKINMTSPYDYHITDDINEHPYSNEEVMLSQVVLNGLYG-----IPAL 511
 G K I+ + H + L+K++ + LYG +P L
 Sbjct: 353 KTGLFKEFIDKWTYVKTTH-----ERGAQKLAKLMFDSLYGKFASNPVDTGKVPYL 403

Query: 512 RSHFNL-FRLDDNNELYNIINGYKNTERNILFSTFVTSRSLYNLLVPPQYLTESEIDNFI 570
 + +L FR+ D YK+ + F+T+ + + + Q D
 Sbjct: 404 KEDGSLGFRVGDEE-----YKDPVYTPM-GVFITAWARFTTITAAQACY-----DRI 449

Query: 571 IYCDTDSLYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHK-----YAYEVNG 625
 IYCDTDS+++ P + + DP LG W E+ + L K YA EV+G
 Sbjct: 450 IYCDTDSIHLTGTEVPEIHKDIDVPKKGWYAHES-TFKRAKYLRQKTYIQDIYAKEVDG 508

Query: 626 KI 627
 K+
 Sbjct: 509 KL 510

>gi|1572479|emb|CAA65712| (X96987) DNA polymerase [Bacteriophage
 GA-1]
 Length = 578

Score = 46.1 bits (107), Expect = 0.001
 Identities = 80/376 (21%), Positives = 146/376 (38%), Gaps = 54/376 (14%)

Query: 234 QLYTYIHNDVILGMCHIHYSDFPNFDYNKLTFSLNIMESYLNEMTRFQLLNQYQDIKI 293
 ++ Y+ +D++I+ + +F N D+ +T + + +Y EM + +Y +
 Sbjct: 162 EIEYLKHDLLIVALA---LRSMFDN-DFTSMTVGSALNTY--KEMLGVKQNEKYFPVL- 214

Query: 294 SYTHYHFDNMNFYDIKSFYRGGLNMYNTKYINKLIDEPFCSIDINSSYPYVMYHEKIPT 353
 + I+ Y+GG N KY + + D+NS YP +M ++ +P
 Sbjct: 215 -----SLKVNSEIRKAYKGGFTWVNPKYQGETVYGGMV-FDVNSMYPAMMKNKLLP- 264

Query: 354 WLYFYEHYSEPTLIPTFLDDNYFSLYKIDKDVFNDDLLIKISRVLRQMXXXXXXXXXX 413
 Y EP + + + LY F + KI ++

286

Sbjct: 265 -----YGEPMVFKGEYKQVVEYPLYIQVRCFFELKKDKIPCIQIKGNARFGQNEYLS 317
 Query: 414 XXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLNKINM 473
 L +T +D I+ + + I+B E+ +P+ + I
 Sbjct: 318 TSGDEYVDLY----VTNVDWELIKKH-YDIFEEFIGG--FMFKGF-----IGF 359
 Query: 474 TSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYGIPALRSHFN--LFRLLDNNELYNIIN 531
 Y + N S E+ + +K++LN LYG A + LD+N L
 Sbjct: 360 FDEYIDRFMEIKNSPDSSAEQSLQAKMLNSLYGKFATNPDPITGKVPYLDENGVLKFRKG 419
 Query: 532 GYKNTERNILFST---FVTSRSLYNLLVPFQYLTESEIDNFIYCDTDSLYMKSUVKPLL 588
 K ER+ +++ F+T+ + N+L Q L FIY DTDS++++ + +
 Sbjct: 420 ELK--ERDPVYTPMGCFITAYARENILSNAQKLYP-----RFIYADTDSIHVEGLGEVDA 472
 Query: 589 NPSLFDPIALGKWDIE 604
 + DP LG WD E
 Sbjct: 473 IKDVIDPKLGYWDHE 488

>gi|118851|sp|P06950|DPOL BPPZA DNA POLYMERASE (EARLY PROTEIN GP2)
 >gi|75812|pir|ERBP2Z DNA-directed DNA polymerase (EC
 2.7.7.7) - phage PZA >gi|216051 (M11813) gene 2 product
 [Bacteriophage PZA] >gi|224741|prf|1112171E ORF 2
 [Bacteriophage PZA]
 Length = 572

Score = 45.3 bits (105), Expect = 0.002
 Identities = 98/461 (21%), Positives = 166/461 (35%), Gaps = 110/461 (23%)

Query: 198 QLKTDNFYITFDKNDMDNDSEAYDYAVKCFAKLTPEQLTYIHNDVIIIGMCHIHYSIDIFP 257
 ++ DF T+ D D + Y ++TP++ YI ND+ I+ + I
 Sbjct: 129 KIAKDFKLTVLKGDIDYHKERPVGY-----EITPDEYAYIKNDIQIIEALL----IQF 178
 Query: 258 NFDYNKLTFSNLNIMESYLNEMTR-----FQLLNQYQDIKISYTHYHFDMNFYDIKSF 312
 +++T + + + + + T+ F L+ D ++ Y
 Sbjct: 179 KQGLDRMTAGSDDLKGFKDIIITKKFKVPTLSLGLDKEVRYA-----222
 Query: 313 YRGGNMYNTKYINKLIDPCFSIDINSSYPYVMYHEKIPTWLYFYEHYSEPTLIPT--F 370
 YRGG N ++ K I E D+NS YP MY +P Y EP +
 Sbjct: 223 YRGGFTWLNDRPFKEKEIGEGMV-FDVNSLYPAQMYRLLP-----YGEPIVFEQYV 273
 Query: 371 LDDDNYSFLYKID-----KDVFNDDLIIKISRVLRQMXXXXXXXXXXXXXXXXXXLRMI 425
 D+D + I K+ + + IK +SR +
 Sbjct: 274 WDEYDPLHIQHRCFELKEGYIPTIQIK-RSRFYKGNEYLKSSGGEIADLW-----324
 Query: 426 QDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIKTQGKLNKINMTSPYDYHITDDI 485
 ++ +D + + + +Y EY F T G K+ I+ + I
 Sbjct: 325 --VSNDV-LELMKEHYDLYNVEYISGLK-----FKATTGLFKDFIDKWTHIKTTSEGA 375
 Query: 486 NEHPYSNEEVMLSKVVLNGLYG-----IPALRSHFN--LFRLLDNNELYNIINGY 533
 + L+K++LN LYG +P L+ + L FRL G
 Sbjct: 376 KQ-----LAKMLNSLYGKFASNPDPVTGKVPYKENGALGFR-----GE 415
 Query: 534 KNTERNIL--FSTFVTSRSLYNLLVPFQYLTESEIDNFIYCDTDSLYMKSUVKPLLNPS 591
 + T+ + F+T+ + Y + Q D IYCDTDS+++ P +
 Sbjct: 416 EETKDPVYTPMGVFITAWARYTTITAAQACF-----DRIIYCDTDSIHLTGTEIPDVIK 470
 Query: 592 LFDPIALGKWDIENEQIDKMFVLNKKYAY-----EVNGKI 627
 + DP LG W E+ + L K Y EV+GK+
 Sbjct: 471 IVDPKLGYWAHES-TFKRAKYLRQKTYIQDIYMKEVDGKL 510

>gi|2435429 (AF012250) unassigned reading frame (possible DNA
 polymerase) [Physarum polycephalum]
 Length = 544

Score = 44.9 bits (104), Expect = 0.002
 Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)

Query: 179 TSIATLGKLLDGGYLTESQLKTDNFYITFDKNDMDNDSEAYDYAVKCFAKLTPEQLTYI 238
 T + L K L D + T Q F N M Y + CF L P++ I
 Sbjct: 62 TQLENLLKSLQDSSFYTFKQ-----FTYQNM-----YSLBISCF--LYPKKKILI 105
 Query: 239 HNDVIIIGMCHIHYSIDIFPNFD-----YNKL--TFSNLIMESY-LNNEMTRFQLLNQYQD 290
 D+ +I Y+D+ ++ YN++ +++NI Y L+ ++ +
 Sbjct: 106 -KDLNFFSENIYNDVVKDYKLLAILYNEIQAYNININRKYILSTASLSLRFKKSFP 164

Query: 291 IKISYTHYHFDNMNFYDIKSFYRGGLNMYNTKYINKLIDEPFCFSIDINSSYPYVMYHEK 350
 K + D + +YI+ Y GG N I + + + + D+NS YPY+M EK
 Sbjct: 165 EKYLRIPLHTRDED--NYIRKSYIGGRNE-----IFEHVAQRNYFYDVNSLYPYIMKKEK 217

Query: 351 IPTWLYFYEHYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDLL---IKIKSRVLRQ 402
 +P + Y + + F + +N+F L I+K N +L + IK+ V
 Sbjct: 218 MPIGI---PEYRDKEYMKKFEKNIENFFGFDVLITIEKTNNNIPVLPYRMGIKNNV-EV 273

Query: 403 MXXXXXXXXXXXXXXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
 L + Q I+ IY + ++++F+ Y +
 Sbjct: 274 GIIYAKGTLRGIYFSEEIKLALKQGYKIE-----IYSAYEYKEKEVVFEEYVEQ 323

Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPALRS 513
 + LK K D + D L K +LN LYG I +
 Sbjct: 324 MYNRRLKAK-----DPALKD-----LYKKLLNTLYGRGLVYEQIDIISP 363

Query: 514 HFNLFRLDDNNELNYIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDNFIYC 573
 L + DN + + + + N ++ + + + F Y T + + IY
 Sbjct: 364 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNLYNLHVIYI 421

Query: 574 DTDLSYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYAY-EVNGKIKIASA 632
 DTD L++K+ P+ + +L +GK+ +E+ + F+ N K Y Y +N I
 Sbjct: 422 DTGGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 477

Query: 633 GIPK-----NAFDTSVDFETFVR----EQFFDGAIENNKSIYNEQGT-----ISIIYPSK 678
 GIP N D + + +F +I NN Y+ Q + I Y +
 Sbjct: 478 GIPLQKPIFNHDIITQHKKILNITLGHYFTFSIRLNNNQYTSFQASRKRLIPNYKTT 537

Query: 679 TEIVC 683
 I+C
 Sbjct: 538 PWIIC 542

>gi|1084487|pir||S41618 DNA polymerase - slime mold (Physarum
 polycephalum) >gi|509721|dbj|BAA06121.1| (D29637) DNA
 polymerase [Physarum polycephalum]
 Length = 547

Score = 44.9 bits (104), Expect = 0.002
 Identities = 118/545 (21%), Positives = 206/545 (37%), Gaps = 104/545 (19%)

Query: 179 TSIATLGKLLDGGYLTESQLKTDFTNYTIFDKDNDMNDSEAYDYAVKCFAKLTPEQLTYI 238
 T + L K L D + T Q F N M Y + CF L P++ I
 Sbjct: 65 TQLFNLKSLQDSSFYTFKQ-----FTYQNM-----YSLEISCF--LYPKKKILI 108

Query: 239 HNDVILGMCHIHYSIDIFPNFD-----YNKL--TFSLNIMESY-LNNEMTRFQLLNQYQD 290
 D+ +I Y+D+ ++ YN++ +++NI Y L+ ++ +
 Sbjct: 109 -KDLNPFSENIYNDVVKDYKLLAILYNEIQTAYNININRKYILSTASLSLRIFKKSFP 167

Query: 291 IKISYTHYHFDNMNFYDIKSFYRGGLNMYNTKYINKLIDEPFCFSIDINSSYPYVMYHEK 350
 K + D + +YI+ Y GG N I + + + + D+NS YPY+M EK
 Sbjct: 168 EKYLRIPLHTRDED--NYIRKSYIGGRNE-----IFEHVAQRNYFYDVNSLYPYIMKKEK 220

Query: 351 IPTWLYFYEHYSEPTLIPTFLDD-DNYFS----LYKIDKDVFNDDLL---IKIKSRVLRQ 402
 +P + Y + + F + +N+F L I+K N +L + IK+ V
 Sbjct: 221 MPIGI---PEYRDKEYMKKFEKNIENFFGFDVLITIEKTNNNIPVLPYRMGIKNNV-EV 276

Query: 403 MXXXXXXXXXXXXXXXXXXXXLRMIQDITGIDCMHIRVNSFVIYECEYFHARDIIFQNYFIK 462
 L + Q I+ IY + ++++F+ Y +
 Sbjct: 277 GIIYAKGTLRGIYFSEEIKLALKQGYKIE-----IYSAYEYKEKEVVFEEYVEQ 326

Query: 463 TQGK-LKNKINMTSPYDYHITDDINEHPYSNEEVMLSKVVLNGLYG-----IPALRS 513
 + LK K D + D L K +LN LYG I +
 Sbjct: 327 MYNRRLKAK-----DPALKD-----LYKKLLNTLYGRGLVYEQIDIISP 366

Query: 514 HFNLFRLDDNNELNYIINGYKNTERNILFSTFVTSRSLYNLLVPFQYLTESEIDNFIYC 573
 L + DN + + + + N ++ + + + F Y T + + IY
 Sbjct: 367 EKEL--ITDNTYISHDTTEFIDITANTCYNNIAITSAITSYARIFMYNTILNLYNLHVIYI 424

Query: 574 DTDLSYMKSVVKPLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYAY-EVNGKIKIASA 632
 DTD L++K+ P+ + +L +GK+ +E+ + F+ N K Y Y +N I
 Sbjct: 425 DTGGLFLKN---PIPDIALTTSKEMGKFRLESINAEAHFIAN-KFYIYAPINSPIIYKFK 480

Query: 633 GIPK-----NAFDTSVDFETFVR----EQFFDGAIENNKSIYNEQGT-----ISIIYPSK 678

288

GIP N D + + +F +I NN Y+ Q + I Y +
 Sbjct: 481 GIPLQKPIFNHDIITQHKKILNITLGHYFTFSIRLNNNQTYSFQASRKRKLIPNYKTT 540

Query: 679 TEIVC 683

I+C

Sbjct: 541 PWIIC 545

>gi|4877819|gb|AAD31446.1| (AF133505) DNA polymerase [Neurospora crassa]
 Length = 1035

Score = 44.1 bits (102), Expect = 0.004
 Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)

Query: 521 DDNNELYNIIINGYKNTERNILFSTFVTSRSLYNLLVFPQYLTESEIDDNFIYCDTDSLIM 580
 + N EL + ++G K+ I ++ + + + + + + S Y DTDS+++

Sbjct: 817 EKNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA-----YTDTSIFV 870

Query: 581 KSVVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYYAYEVNGKIKIASAGIPKNAFD 640
 + KPL + + + K + + I + ++ K Y + GK++I GI KN +

Sbjct: 871 E---KPLDSAFIGEGCGKFAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 927

Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
 T+ + + E ++G + + E GT+++ K ++ G YD+

Sbjct: 928 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 977

>gi|461962|sp|P33537|DPOM_NEUCR PROBABLE DNA POLYMERASE
 >gi|283351|pir|S26985 probable DNA-directed DNA
 polymerase (EC 2.7.7.7) - Neurospora crassa
 mitochondrion plasmid maranhar (SGC3)
 >gi|578156|emb|CAA39046| (X55361) putative DNA
 polymerase [Neurospora crassa]
 Length = 1021

Score = 44.1 bits (102), Expect = 0.004
 Identities = 36/172 (20%), Positives = 82/172 (46%), Gaps = 14/172 (8%)

Query: 521 DDNNELYNIIINGYKNTERNILFSTFVTSRSLYNLLVFPQYLTESEIDDNFIYCDTDSLIM 580
 + N EL + ++G K+ I ++ + + + + + + S Y DTDS+++

Sbjct: 815 EKNYELLSYLDGEKDDGFIINSTSIAAATASWSRILMYKHIINSA-----YTDTSIFV 868

Query: 581 KSVVVKPLLNPSLFDPIALGKWDIENEQIDKMFVLNHHKYYAYEVNGKIKIASAGIPKNAFD 640
 + KPL + + + K + + I + ++ K Y + GK++I GI KN +

Sbjct: 869 E---KPLDSAFIGEGCGKFAEYNGQLIKRAIFISGKLYLLDFGGKLEIKCKGITKNKDN 925

Query: 641 TSVDFETFVREQFFDG---AIIENNKSIYNEQGTISIYPSKTEIVCGNVYDE 689
 T+ + + E ++G + + E GT+++ K ++ G YD+

Sbjct: 926 TTHNLDINDFEALYNGESRVLFQERWGRSLELGTVTVKYQKYNLISG--YDK 975

>gi|2499511|sp|Q12471|6P22_YEAST 6-PHOSPHOFRUCTO-2-KINASE 2
 (PHOSPHOFRUCTOKINASE 2 II) (6PF-2-K 2)
 >gi|2131162|pir|S61066 6-phosphofructo-2-kinase (EC
 2.7.1.105) - yeast (Saccharomyces cerevisiae)
 >gi|2131163|pir|S71026 6-phosphofructo-2-kinase (EC
 2.7.1.105) - yeast (Saccharomyces cerevisiae)
 >gi|1085116|emb|CAA62371| (X90861)
 6-phosphofructo-2-kinase [Saccharomyces cerevisiae]
 >gi|1420028|emb|CAA99157| (Z74878) ORF YOL136c
 [Saccharomyces cerevisiae] >gi|1628439|emb|CAA64733|
 (X95465) 6-phosphofructo-2-kinase [Saccharomyces
 cerevisiae]
 Length = 397

Score = 40.6 bits (93), Expect = 0.041
 Identities = 48/208 (23%), Positives = 92/208 (44%), Gaps = 29/208 (13%)

Query: 175 MKTNTSIATLGKLLDGGYLTESQLKTDNFNYTIFDKNDMNDSEAYDYAVKCFKLTPEQ 234
 ++ S AT+ K LL L+ + + FN K+ND ++ +A++T ++

Sbjct: 139 IRRQISCATISKPLL----LSNTSSDLFN----PKNNDKKET-----YARITLQK 181

Query: 235 LTY-IHNDVILGMCHIHYSDFPNFDYNTLFTSLNIMESYLNEMTRFOLLN---QYQD 290
 L + I+ND +G+ S I + F + S+ +E++ F L+ Q

Sbjct: 182 LFHEINNDECDVGIFDATNSTI-----ERRRFEEVCSFNTDELSSFNLVPIILQVSC 235

289

Query: 291 IKISYTHYHFHDMNFY-DYIKSFYRGGLNMYNTKYINKLIDEPFCFSID-INSSYPYVMYH 348
 S+ Y+ H+ +F DY+ Y + + + + FS+D N + Y+ H
 Sbjct: 236 FNRSEIKYNIHNKSFNEDYLDKPYELAIAKDFAKRLKHYYSQTFPSLDEFNQIHRYISQH 295

Query: 349 EKIPTWLYFYEYHEPTLIPTFLDDNY 376
 E+I T L+F+ + + P L+ +Y
 Sbjct: 296 EEIDTSLFFFNVINAGVVEPHSLNQSHY 323

>gi|2258375|gb|AAD11909.1| (AF007261) transcription initiation
 factor sigma [Reclinomonas americana]
 Length = 532

Score = 39.9 bits (91), Expect = 0.070
 Identities = 49/205 (23%), Positives = 84/205 (40%), Gaps = 14/205 (6%)

Query: 100 NHFLLKDTMRYFDNITRENIYLKSAEENEHTLKMKEATILAKNQNVIL---EKRVKSSIN 156
 N+ + + F + ++IY+ + +KE L K NVI+ K +K N
 Sbjct: 177 NYLVKNSYLNLFKTVPHDSIYMNSYIQTPLNILKEYLQLIKIINVIILQINKNIKKNN 236

Query: 157 LDLMFLNGFKFNIIDNFM---KTNTSIATLGKKLLDGGYLTESQLKTDNFYTFDKDND 213
 L+++FL F + N++ K + + + K L Y+T L T Y K
 Sbjct: 237 LNISLFLYKIFYQELKWNIFYINKISRNTQKINIKTLKNSYITFYNLITFIQYTTKKQRL 296

Query: 214 MNDSEAYDYAVKCFK--LTPEQLTYIHNDVIIIGMCHIHYSDFPNFDYN-KLTFSINI 270
 D +K F K P+ +N +I G+ HI+ + N K+T I
 Sbjct: 297 KKDIFYKQIFIKTFLKQHKIPKINKKNSLIKYGLTHIYDMILISILRENKVTLKQRI 356

Query: 271 MESYLNEMTRFQLLNQYQDIKISY 295
 + +Y+ T + QY +KI Y
 Sbjct: 357 IFNYMPYITT---ISKQY--VKIGY 376

>gi|15734|emb|CAA37450| (X53370) DNA polymerase (AA 1-575)
 [Bacteriophage phi-29]
 Length = 575

Score = 39.5 bits (90), Expect = 0.092
 Identities = 41/150 (27%), Positives = 64/150 (42%), Gaps = 36/150 (24%)

Query: 497 LSKVVLNGLYG-----IPALRSHFNL-FRLDDNNELNYNIINGYKNTERNIL--F 542
 L+K++LN LYG +P L+ + L FRL G + T+ +
 Sbjct: 381 LAKLMLNSLYGKFASNPVDVTGKVPYKENGALGFRL-----GEEETKDPVYTPM 429

Query: 543 STFVTSRSLYNLLVPFQYLTESEIDNFIYCDTDSLVMKSVVKKPLNPSLFDPIALGKWD 602
 F+T+ + Y + Q D IYCDTDS+++ P + + DP LG W
 Sbjct: 430 GVFITAWARYTTITAAQACY-----DRIIYCDTDSIHLTGTEIPDVIKDIVDPKKLGWYA 484

Query: 603 IENEQIDKMFVLNHHKYAY----EVNGKI 627
 E+ ++ L K Y EV+GK+
 Sbjct: 485 HES-TFKRVKYLKQKTYIQDIYMKEVDGKL 513

Query= pt|110872 44AHJDORF002 Phage 44AHJD ORF |3789-5732|3 1
 (647 letters)

>gi|135273|sp|P27622|TAGC_BACSU TEICHOIC ACID BIOSYNTHESIS PROTEIN C
 >gi|478126|pir|D49757 teichoic acid biosynthesis protein
 tagC - Bacillus subtilis (strain 168) >gi|143727
 (M57497) putative [Bacillus subtilis]
 >gi|2636103|emb|CAB15594.1| (Z99122) alternate gene
 name: dinC [Bacillus subtilis]
 Length = 442

Score = 112 bits (278), Expect = 7e-24
 Identities = 91/314 (28%), Positives = 147/314 (45%), Gaps = 58/314 (18%)

Query: 152 FELNELEPKFVMGFGGIRNAVNSINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
 F+ + PK V QS N D++ + +Y+TQ S + + I +L+ G
 Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLGKDNTQSYRITRLSLEG 66

Query: 208 DLISSMRIVQGGHGTIGLERQSNNGEMKIWLHHD-----GVAKLLQVAYKDNVYLDLEEA 262
 + SM + GGHGT IG+E + NG + IW +D ++L+ YK LD E +
 Sbjct: 67 LQLDSMLLKHHGGHTNIGIENR-NGTIYIWSLYDKPNETDKSELVCFPYKAGATLD-ENS 124

290

Query: 263 KGLTDYTPQSLNKHFTFTPLIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDN 322
 K L ++ H TP +D N +L +R + D KN+ N ++ +TI N
 Sbjct: 125 KELQRFSNMPP--DHRVTPALDMKNRQLAIR-----QYDTKNN--NNKQWVTIFN 170

Query: 323 SE-----NNDN-----RWMQGIADVGDDBLYWLSGNSSVNSHVQIGKYSLTGTGQKI 367
 + N +N ++QG +D LYW +G+++ S+ + +
 Sbjct: 171 LDDAIANKNNPLYTINIPDELHYLQGFFLDDGYLYWYTGDTNSKSYPNL-----ITV 222

Query: 368 YDYPFKLSYQDGINFPRD-----NFKEPEGICIYTNPKTKRKSLLAMTNGGGGKRFB 420
 +D K+ Q I +D NF+EPEGIC+YTNP+T KSL++ +T+G G R
 Sbjct: 223 FDSDNKIVLQKEITVGKDLSTRYENNFREPEGICMYTNPETGAKSLMVGITSGKEGNRIS 282

Query: 421 NLYGFFQLGEYEHF 434
 +Y + YE+P
 Sbjct: 283 RIYAYH---SYENF 293

>gi|142847 (M64050) DNase inhibitor [Bacillus subtilis]
 Length = 125

Score = 51.9 bits (122), Expect = 1e-05
 Identities = 35/116 (30%), Positives = 55/116 (47%), Gaps = 10/116 (8%)

Query: 152 FELNELEPKFVMGFGGIRNAVNSINIDKETNHMYSTQSDS----QKPEGFWINKLTPSG 207
 F+ + PK V QS N D++ + +Y+TQ S + + I +L+ G
 Sbjct: 7 FDFTNITPKLFTELRVADKTVLQSFNFDEKNHQIYTTQVASGLKNDTQSYRITRLSLEG 66

Query: 208 DLISSMRIVQGGHGTITIGLERQSNEMKIWLHHD-----GVAKLLQVAYKDNVLD 258
 + SM + GGHGT IG+E + NG + IW +D ++L+ YK LD
 Sbjct: 67 LQDLSMLLKHGHTNIGMENR-NGTIYIWSLYDKPNETDKSELVCFYKAGATLD 121

>gi|4038407 (AF103943) factor C protein precursor [Streptomyces
 griseus]
 Length = 324

Score = 39.1 bits (89), Expect = 0.10
 Identities = 61/269 (22%), Positives = 102/269 (37%), Gaps = 33/269 (12%)

Query: 172 VNQSINIDKETNHMYSTQSDSQKPEG---FWINKLTPSGDLISSMRIVQGGHGTITIGLER 228
 V QS D ++ Q S P+ I +L SG+ + M ++ GHG +IG +
 Sbjct: 66 VQQSFTFDIVNRRFLVFAQLKSGSPDDSGDLCTQLDFSGNKLGHMYLLGFHGVSIGAQ- 124

Query: 229 QSNEMKIWLHHDGVAKLLQVAYKDNVLDLEEAKGLTDYTPQSLNKHFTFTP----- 281
 + +W D + + + + G T S L KH P
 Sbjct: 125 PVGADTYLWTEVD-----VNSNARGTRLARFKWNGATLSRTSSALAKHQVPVPGATEMTC 179

Query: 282 LIDEANDKLILRFGDGTIQVRSRADVKNHIDNVEKEMTIDNSENNDNRWMQGIADVGDDBL 341
 ID N+++ +R+ + + +V + V + D QG A+ G +
 Sbjct: 180 AIDPVNNRMAIRYLTASGRRYGIYNVADIAAGVYDKPLSDVPHPTGLGTFQGYALYGSYV 239

Query: 342 YWLSGN-----SSVNSHVQIGKYSLTGTGQKIYDYPFKLSYQDGINFPRDNFKEPEGIC 394
 Y L+GN + NS+V + TG + + + G F+EPEG+
 Sbjct: 240 YQLTGNPYGPDNPNPGNSYVS--SVDVNTGALVQ----RAFTRAGSTL---TFREPEGMG 290

Query: 395 IYTNPKTKRKSLLAMTNGGGGKRFBHLY 423
 IY + + L L +G G R NL+
 Sbjct: 291 IYRTAAGEVR-LFLGFASGVAGDRRSNLF 318

Query= pt|110873 44AHJDORF003 Phage 44AHJD ORF |6626-8389|2 1
 (587 letters)

>gi|138123|sp|P04331|VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9)
 >gi|75850|pir|WMBPT9 gene 9 protein - phage phi-29
 >gi|215327 (M14782) tail protein [Bacteriophage phi-29]
 >gi|225364|prf||1301270D gene 9 [Bacillus sp.]
 Length = 599

Score = 92.4 bits (226), Expect = 8e-18
 Identities = 126/618 (20%), Positives = 251/618 (40%), Gaps = 71/618 (11%)

Query: 5 TNFKFFYNTPTFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPY-NFIRDMEINV 62
 TN + + PF+ DY+NT F S+ + ++F R + + SK + F ++ ++V
 Sbjct: 9 TNVRILADVPFSDNYKNTRWFTSSSNQYNWF--NRKSRVYEMSKVTFMGFRENKPYVSVS 66

Query: 63 MQWHAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
 + Y+ F + D+ ++ +YAFV ++E+ N V ++F ID + T+ ++
 Sbjct: 67 LPIDKLYSASYIMFQADYGNKWFYAFVTELEFKNSAVTYVHFEIDVLQTMFDMKFQES 126

Query: 122 SNVNIERQHLSKRTYNYMLPMLRNDDVLKVSNNKYNVYNQMQQYLENLVLFQSSADLSK 181
 I R+H+ K + P + D+ L ++ + + + ++F S
 Sbjct: 127 F---IVREHV-KLWDDGTPTINTIDEGLSYGSEYDIVSVENHKPYDDMMFLVIIKSKSIM 182

Query: 182 FGT--KKEPNLDTSGGTIYDNITSPVNLVMEYGDFFINMDKMSAYPWITQNFQK----V 235
 GT ++E L+ ++ + + P+ Y+ + + D +I N V
 Sbjct: 183 HGTGPEEESRLNDINASL-NGMPQPLCYIHPF-----YKDGKVPKTYIGDNNANLSPIV 236

Query: 236 QMLPKDFINTKDLVDKTSKITGLKTLKQGGKSKEWSLK-DLSL-----SFSNLQ 285
 ML F + D+ + +T LK K+ + LK D + N+
 Sbjct: 237 NMLTNIFSQSAVNDI-VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVD 295

Query: 286 EMMLSK-----KDEFKHMIRNEYMTIEFYDWNNGNTMLLDAGKISQK 326
 + + K KD+ ++ Y E D+ GN M L I+
 Sbjct: 296 TIFVKKIPDYEALEIDTGDKGWGGFTKQESKLMMPYCVTEITDFKGNHMLNKTEYINNS 355

Query: 327 TGVKLRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQV 386
 +K++ + +G N+V DYN+ D + N+ S +N N
 Sbjct: 356 K-LKIQVRGSLGVSNNKVAYSVDYNA---DSALSGGNRLTASLDSSLINNPN----- 404

Query: 387 PILINNGLQSQANRQ--KNAESQLITNRIDNVLNG---SDPKSRFYDAVSVAANLSP 441
 I I N L Q N+ +N +S ++ N I ++ G + + A+ +AS++
 Sbjct: 405 DIAILNDYLSAYLQGNKNSLENQKSSILFNGIMGIGGGISAGASAAGGSALGMASV-- 462

Query: 442 TALFGKFNEEYNYFYKQQAQYKDLALQPPSVTESEMGNFQIANSINGLTMKISVPSPKE 501
 T + + QA+ D+A PP +T+ AF N G+ + +
 Sbjct: 463 TGMTSTAGNAVLQMAMQAKQADIANIPPQLTKMGNTAFDYGNVGRGVYVIKKQLKAEY 522

Query: 502 ITFLQKYMLFGFEVNDYNSFIEPINSMTVCNLYKCTGTYYTIRDIDPMLMEQLKAILESG 561
 L ++ +G+++N + + NY++ + DI+ +++++ I ++G
 Sbjct: 523 RRLSLSSFHKYGYKINRVKK--PNLRTKAFNYVQTKDCFISGDINNNDLQEIIRTIFDNG 580

Query: 562 VRFWHDGSGNPMQLQNPL 579
 + WH D GN ++N L
 Sbjct: 581 ITLWHTDNIGNYSVENEL 598

>gi|138124|sp|P07534|VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)
 >gi|75849|pir|WMBP9Z gene 9 protein - phage PZA
 >gi|216058 (M1813) tail protein (Bacteriophage PZA)
 Length = 599

Score = 81.9 bits (199), Expect = 1e-14

Identities = 127/618 (20%), Positives = 248/618 (39%), Gaps = 71/618 (11%)

Query: 5 TNKFFFYNTPFT-DYQNTIHFNSNKERDDYFLNGRHFKSLDYSKQPYNFIRDME-INVD 62
 + + PF+ DY+NT F S+ + ++F + + SK + R+ I+V
 Sbjct: 9 TNVRILADVFPFSNDYKNTRWFTSSSNQYNWF--NSKTRVYEMSKVTFQGFRENKSYISVS 66

Query: 63 MQWHAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTYTQGNVLEQL 121
 ++ Y+ F + D+ ++ +YAFV ++EY N ++F ID + T+ N+ Q
 Sbjct: 67 LRLDLLYNASYIMFQADYGNKWFYAFVTELEKYNVGTITYVHFEIDVLQTW-MFNIKQFE 125

Query: 122 SNVNIERQHLSKRTYNYMLPMLRNDDVLKVSNNKYNVYN--QMQQYLENLVLFQSSADLS 179
 S I R+H+ K + P + D+ L ++ + + + Y + + L S +
 Sbjct: 126 SF--IVREHV-KLWDDGTPTINTIDEGLNYGSEYDIVSVENHRPYDDMMFLVVISKSIM 182

Query: 180 KKEGTTKEPNLDTSGGTIYDNITSPVNLVMEY-----GD-----FINMDK 221
 + E L+ ++ + + P+ Y+ + GD +N +
 Sbjct: 183 HGTAGEAESRLNDINASL-NGMPQPLCYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTN 241

Query: 222 MSAYPWITQNFQKQVQMLPKDFINTK-----DLEDVKTSEKITGLKTLKQGGKSKEWS 273
 + + N V M D+I K +L+ K + G+ K G +
 Sbjct: 242 IFSQKSAVNNI--VNMYVTDYIGLKLDYKNGDKELKLDKDMFEQAGIADDKHGNVDITIFV 299

Query: 274 LKDL---SLSFSNLQEMMLSKKDEFKHMIRNEYMTIEFYDWNNGNTMLLDAGKISQKTGVK 330
 K +L + KD+ ++ Y E D+ GN M L I +K
 Sbjct: 300 KKIPDYETLEIDTGDKGWGGFTKQESKLMMPYCVTEVTDFKGNHMLNKTEYIDNNK-LK 358

Query: 331 LRTKSIIGYHNEVRVYPVDYNSAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILI 390
 ++ + +G N+V DYN+ + L+ + L+T++ N+ + I+

292

Sbjct: 359 IQVRGSLGVSNNKVAISIQDYNAGGS----LSGGDRLTAS----LDTSLINNNPNDIAII- 409

Query: 391 NNGILGQSQQANRQ--KNAESQLITNRIDNVNLSGDPKSRFYDAVSASNLSP----- 441
 N L Q N+ +N +S ++ N I +L G A + A SP

Sbjct: 410 -NDYLSAYLQGNKNSLENQKSSILFNGIVGMLGG-----VSAGASAVGRSPFGLASSV 462

Query: 442 TALFGKFNEEYNYFYKQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKISVPSPKE 501
 T + + QA+ D+A PP +T+ AF N G+ + +

Sbjct: 463 TGMTSTAGNAVLDLQALQAKQADIANIPQLTKMGNTAFDYGNGYRGVYVIKQLKAEY 522

Query: 502 ITFLQKYMLFGFEVNDYNSFIEPINSMTVCNYLKTGTGTYTIRDIDPMLMEQLKALESG 561
 L ++ +G+++N + + NY++ + DI+ +++++ I ++G

Sbjct: 523 RRLSLSSFFHKYGYKINRVKK--PNLRTKAYNYIQTKDCFISGDINNNDLQEIRTIFDNG 580

Query: 562 VRFVHNDGSGNPMLQNPL 579
 + WH D GN ++N L

Sbjct: 581 ITLWHTDDIGNYSVENEL 598

>gi|1429238|emb|CAA67657| (X99260) tail protein [Bacteriophage B103]
 Length = 598

Score = 77.6 bits (188), Expect = 2e-13
 Identities = 130/623 (20%), Positives = 240/623 (37%), Gaps = 86/623 (13%)

Query: 5 TNFKFFYNTPT-DYQNTIHFSNKRDDYFLNGRHFKSLDYSKQPYNFI---RDRMEIN 60
 T+ + F N PF+ DY++T F + + YF + K + NF+ I

Sbjct: 9 TDVRIFSNVFNSNDYKSTRWFTNADAQYSYF---NAKPRVHVINECNFVGLKEGTPHIR 64

Query: 61 VDMQWHDAAQGINYMTFLS-DFEDRRYYAFVNQIEYVNDVVVKIYFVIDTITMTYTQGNVLE 119
 V+ + D YM F + + ++ +Y FV ++EYVN V +YF ID I T+ +

Sbjct: 65 VNKRIDDLNACYMIFRNTQYSNKNWFYCFVTRLEYVNSGVNTLYFEIDVIQIW-MFDFKF 123

Query: 120 QLSNVNIERQHLSKRTYNYMLPMLRNDDVLKVSNNKYNVYNQMQQYLENLVLFQSSADLS 179
 Q S + E Q + P+ D+ L + V Q ++F S

Sbjct: 124 QPSYIVREHQMWDANNE---PLTNTIDEGLNYGTEYDVVAVEQYKPYGDLFMVCISKS 180

Query: 180 KKFGTKEPNLDTSGKTIYDNITS---PVNLYVMEYGDFINFMDKMSAYPWITQNFQKVQ 236
 K T E G I NI P++ YV + + D S P +T +VQ

Sbjct: 181 KMHATAGET---FKAGEIAANINGAPQLSYVHPF-----YEDGSS--PKVTIGSNEVQ 230

Query: 237 ML-PKDFINTKDLEDVKTSEKITGLKT-----LKQGGKSKEWSLKDLSLSFSNL----- 284
 + P DF+ ++ + ++ T + +K SL+D + +

Sbjct: 231 VSKPTDFLKNMFTQEHAVNNIVSLYVTDYIGLNIHYDESAKTMSLRDTMFEHAQIADDKH 290

Query: 285 -----QEMMLSKKDEFKHMIRNEYMTIEFY-----DWNNGNTMLLDAGK 322
 +E + +F NE + Y D+ GN + +

Sbjct: 291 PNVNTIYLKEVKEYEKTIDTGKFFASFANNEQSKLLMPYCVTTITDFKGNQIDIKNEY 350

Query: 323 ISQKTGVKLRTKSIIGYHNEVRVYPVDYNS---AENDRPILAKNKEILIDTGSFLNTNIT 379
 ++ + +K++ + +G N+V DYN+ D+ + A NT++

Sbjct: 351 VNG-SNLKIQVRGSLGVSNNKVITYSVQDYNADTTLSGDQNLTA-----CNTSLI 398

Query: 380 FNSFAQVPILINNGILGQSQQANRQ--KNAESQLITNRIDNVNLSGDPKSRFYDAVS 434
 N+ V I+ N L Q N+ +N + ++ N + ++L G+ + AV

Sbjct: 399 NNNPNVDVAII--NDYLSAYLQGNKNSLENQKDSILFNGVMSMLGNGIGAVGSAATGSAVG 456

Query: 435 VASNLSPALFGKFNEEYNYFYKQQAEYKDLALQPPSVTESEMGNAFQIANSINGLTMKI 494
 VAS S T + + QA+ D+A PP + + A+ N G+ +

Sbjct: 457 VAS--SATGMVSSAGNAVLIQGMQAKQADIANTPPQLVKMGNTAYDYGNGYRGVYVIK 514

Query: 495 SVPSPKEITFLQKYMLFGFEVNDYNSFIEPINSMTVCNYLKTGTGTYTIRDIDPMLMEQL 554
 + L + +G++ N + + + NY++ I +++ +++++

Sbjct: 515 KQIKEEYRNILSDFSRKYGYKTNLVK--MPNLRTRESYNYVQTKDCNIIGNLNEDLQKI 572

Query: 555 KALESGVRFVHNDGSGNPMLQN 577
 + I +SG+ WH D G+ L N

Sbjct: 573 RTIFDSGITLWHADPVGDYTLNN 595

>gi|215339 (M12456) p9 tail protein [Bacteriophage phi-29]
 >gi|224163|prf||1011232C protein p9,tail [Bacteriophage
 phi-29]
 Length = 335

293

Score = 71.0 bits (171), Expect = 2e-11
 Identities = 64/293 (21%), Positives = 123/293 (41%), Gaps = 20/293 (6%)

Query: 292 KDEFKHMIRNEYMTIEFYDWNNTMLLDAGKISQKTGVKLRTKSIIGYHNEVRVYPVDYN 351
 KD+ ++ Y E D+ GN M L I+ +K++ + +G N+V DYN
 Sbjct: 57 KQESKLMYPYCVTEITDFKGNHMLKTEYINNSK-LKIQVRGSLGVSNNKVAYSVDYN 115

Query: 352 SAENDRPILAKNKEILIDTGSFLNTNITFNSFAQVPILINNGILGQSQQANRQ--KNAES 409
 + D + N+ S+N N I I N L Q N+ +N +S
 Sbjct: 116 A---DSALSGGNRLTASLDSSLINNNPN-----DIAILNDYLSAYLQGNKNSLENQKS 165

Query: 410 QLITNRIDNVLNG---SDPKSRFYDAVSASNLSPALFGKFNEEYNYFYKQQAEYKDLA 466
 ++ N I ++ G + + A+ +AS++ T + + QA+ D+A
 Sbjct: 166 SILFNGIMGIGGGISAGASAAGGSALGMASV--TGMTSTAGNAVLQMAMQAKQADIA 223

Query: 467 LQPPSVTESEMGNFQIANSINGLTMKISVPSPEITFLQKYMLFGFEVNDYNSFIEPI 526
 PP +T+ AF N G+ + + L ++ +G+++N +
 Sbjct: 224 NIPPQLTKMGGNTAFDYGNGYRGVYVIKKQLKAEYRRSLSSFFHKYGYKINRVKK--PNL 281

Query: 527 NSMTVCNYLKCTGTYTIRIDPMLMEQLKAILESGVRFWHNDGSGNPMLQNPL 579
 + NY++ + DI+ +++++ I ++G+ WH D GN ++N L
 Sbjct: 282 RTRKAFNYVQTKDCPISGDINNNDLQEIRTIFDNGITLWHTDNIGNYSVENEL 334

>gi|1181968|emb|CAA87738.1| (Z47794) tail protein [Bacteriophage
 CP-1]
 Length = 230

Score = 53.9 bits (127), Expect = 3e-06
 Identities = 29/113 (25%), Positives = 54/113 (47%), Gaps = 3/113 (2%)

Query: 1 MRKLTNFKFFYNTPF-TDYQNTIHFNSNKERDDYFLNGRHFSLDYSKQPYNFIRDRMEI 59
 M++ T + +PF DY N I+F + + +D+F + Y + + + I
 Sbjct: 1 MQESTKIWLYAKSPFKNDYANVINFTRESMEDFTTKNPHIEIYVEYDKFQYQTRNGSI 60

Query: 60 NVDMQWHDAAQGINYMTFLSDFEDRRYYAFVNQIEYVNDVVVKIYFVIDTIMTY 112
 V + + + YM F+++ R YYAFV + Y+N+ +I + +D TY
 Sbjct: 61 VVSGRVEKYENVVTYMRFINN--GRYYAFVFDVLYINEDATRIIYEVDVWNTY 111

>gi|1181970|emb|CAA87740.1| (Z47794) tail protein [Bacteriophage
 CP-1]
 Length = 586

Score = 42.2 bits (97), Expect = 0.010
 Identities = 79/381 (20%), Positives = 139/381 (35%), Gaps = 92/381 (24%)

Query: 277 LSLSFSNLQEMMLSK--KDEFK---HMIRNEYMTIEFYDWNNTMLLDAG----KISQKT 327
 L +++ +QE + S KD+ + ++ +E+ IE YD GN+ + I +
 Sbjct: 187 LKAYDQIQEGLRSYMGKDDLEIEVQLNSEPTEIELYDIYGN SYVYQPQYLPTIDEAH 246

Query: 328 GVKLRTKSIIGYHNEVRVYPVDYNSAEN-----DRPIL----- 360
 K+ +G N+V + ++YN+A N D+ IL
 Sbjct: 247 KYKVIVSGSLGDSNQVHINFLEYNNANNVSYADKNILDSLESQDWAHNPEHFKYGLNDV 306

Query: 361 -AKNKEILIDT-GSFLNTNITFNSFAQVPILINNGILGQSQQANRQKNAESQLITNRIDN 418
 K+ IL D S++ ++ Q+ N +L QS + ++ A + +
 Sbjct: 307 TGKSVAILNDAEASYIQSHKNQMEHTQLTFKENRDMKQSVDLNKNQVATANSQASNAQ 366

Query: 419 VLNGSDPKSRFYDAVSASNLSPALFGKF-----NEEYNYFYKQQQ-- 459
 S +++ + S N++ L G F N +YN QQ
 Sbjct: 367 FAVDSANINQWTEGASGILNVAGNLLTGNFGGALGLASGGMKVFANRDYNDKVYQQGF 426

Query: 460 -----AEYKDLALQPPSVTESEMGNFQIANSIN 488
 A DL QP SV + AFQ N +
 Sbjct: 427 TSENNALKSQSNALANMKSIALDQSI RAYNATMADLQNPISVQQIGNDLAFQSGNRLT 486

Query: 489 GLTMKISVPSPEITFLQKYMLFGFEVNDY-NSFIEPINSMTVCNYLKCTGTY--TIRD 545
 + K+S+ + + +Y +G VN + N + + S NY+K T+R
 Sbjct: 487 DVYWKSLAQKEIMGRANEYIKCYGVLVNWFNDALSVMRSRKRFPNYIKMINVNLGTLR- 545

Query: 546 IDPMLMEQLKAILESGVRFWH 566
 + M ++AI +SGVR W+
 Sbjct: 546 ANQSHMNAIQAIQSGVRIWN 566

294

Query= pt|110875 44AHJDORF005 Phage 44AHJD ORF |12643-13890|-1 1
(415 letters)

>gi|3845203 (AE001399) GAF domain protein (cyclic nt signal
transduct.) [Plasmodium falciparum]
Length = 1245

Score = 52.3 bits (123), Expect = 6e-06
Identities = 59/246 (23%), Positives = 105/246 (41%), Gaps = 27/246 (10%)

Query: 174 ESIDRNHGNVDYIGFPMFLGNAVNFSSPILSNLNIYNLLQKHKMTSRLYKNIFLEMR 233
+S D N+ N + + N+V FS+ N IY++L N +YK + E+
Sbjct: 854 DSSDNNNNNNNNNNNNNNNNNNNSVIFST----NEKIYDML----NRDNIYKVKKEIF 904

Query: 234 RNDYVNEKRNTAFNSNDAMTTGEFNEYNLADDNLRNHINQNGDFFYIKTDDKYI-- 291
D + + + +N+ M + N N ++N+ N+ N NGD Y KY
Sbjct: 905 EGDSSIKTMTENKPNLTNKNVMMNDNIDNNNNNNNNNNIDNNNNNGDNIYNDLKKYYLN 964

Query: 292 KVMYNVTTFMNTIIVVPYTKQYEFCTKIR-DIDNHVTYLRDDMFYKENMERYYNPSNLH 350
++N ++ + + + K E K+ I + L +P+K NM + + L+
Sbjct: 965 TSIFNKDLYVKKHFVDIIMNKSLEEIIKMNVIYISERINSL---LPHKGNM---LNDVTKLY 1018

Query: 351 FDNAYSKNYVVDNDRYLYLDMNKIIKFIKHNEMKKNMSEFERKEKIYEDN----YIENTK 406
NAY + N K I F + E K +M F+ +KIY+ N + N K
Sbjct: 1019 MSNAYGEKCFPFN----FPQIKEIIFVNEYEKMDMKYFKMLKKIYKYNLNKIFSNNYK 1073

Query: 407 KYLMKQ 412
+++K+
Sbjct: 1074 FFIKK 1079

>gi|3758843|emb|CAB11128.1| (Z98551) predicted using hexExon;
MAL3P6.23 (PFC0820w), Hypothetical protein, len: 4982 aa
[Plasmodium falciparum]
Length = 4981

Score = 49.2 bits (115), Expect = 5e-05
Identities = 67/287 (23%), Positives = 110/287 (37%), Gaps = 60/287 (20%)

Query: 127 ITDLNSATDLKYHSNFKHYPIIIYDEFLAEDDYLDIDWDKLT----IYESIDRNHGN 182
I D+N + D+ + +++ I YD +++DK++ IY +ID++ N
Sbjct: 3619 IMDINKSKDISKNMEIVQN---IEYD-----NKYDKIRNDMDAIYMAIDKMDN 3664

Query: 183 VDYIGFPMFLGNAVNFSSPILSNLNIYNL----LQKHKMTSRLYKNIFLEMRRNDYV 238
+ I + F L N S +N YNL ++ K N R Y N F +D
Sbjct: 3665 IGIINCMRYFNLYKNYNLSNECNRRE-YNLNELYMEDIKRNMKR-YDNNFNINHYDDNN 3722

Query: 239 NEKRNTAFNSNDAMTTGEFNEYNLADDNLRNHINQNGDFFYIKTDDKYIKVMYNVT 298
N N N+N++ N N ++N N+ N NG F+ D
Sbjct: 3723 NNGGCGFFHVD----- 3771

Query: 299 TFMNTIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYNPSNLHFDNAYSKN 358
K FCTK ++F +N+E N N N Y+ N
Sbjct: 3772 -----KDLFFCTK-----KNIFPCKNIETVCKNEYNKKIYNNYTCN 3807

Query: 359 YVVDNDRYLYLDMNKIIKFIKHNEMKKNMSEFERKEK-IYEDNYIEN 404
V+N + ++IK + + N E+ + EK +Y + EN
Sbjct: 3808 ISVNNTLNLCLNIKELIKLNNNKKKILNYYEYHKVEKLLYRHSFEN 3854

Score = 35.6 bits (80), Expect = 0.70
Identities = 62/290 (21%), Positives = 121/290 (41%), Gaps = 65/290 (22%)

Query: 2 VKQNRDLMVRDYQNAVN--HVRKKIPDKYNQIELVDELMNDDIDYIISISNRSDGKSFNY 59
+K+N ++ +N+N +V++ DK N I D++I+ SN + +SF
Sbjct: 4445 IKRNNINKSNIKRNINKSNVKSNTDKSNVIS-----DFHIT-SNNNITRSFT- 4492

Query: 60 VSPFIYLAIKLDIKFTLLSRHYTLRDAYRDFIEEIIDENPLFKSRVTFRSARDYLAIY 119
A D F LS TL +Y +F ++ I
Sbjct: 4493 -----ATLTDSIFNTLSE--TLNYSYDNFFSNMDN-----IKI 4523

Query: 120 QDKEIGVITDLNSATDLKYHSNFKHYPIIIYDEFL-----ALEDDYLDIDWDKLTIE 174
+ EI ITD++ +YH N+LK + +E++ + +D + DE ++T+ E
Sbjct: 4524 KKNENINITDVDYGNKKEYHENYLVKVKQKVNEEYIEETFKSDKDCSIKDEACTIRTLSE 4583

295

Query: 175 S--IDRNHGNVDYIGFPMFLGNVNFSSPILSNLNIYNLLQKHKN--TSRLYKNIFL 230
 S I N N+D + + + S P N++ N ++K+ +N R+ KN
 Sbjct: 4584 SCNISENISNID-----MDDEDHISFPNGRNVHDNNYMKKNHVNYDKMRVGGKNIP 4634

Query: 231 EMRRNDYVNEKRNTRAFNSNDDAMTTGEFNEYNLADDNLRNHINQNGD 280
 D + +++ + +D M++ ++ E ++ + L + NG+
 Sbjct: 4635 SFTTFDKILDEKKKK---SDKDMSSSKWLEREEHIKEIKLEKNYMNMG 4680

Score = 34.0 bits (76), Expect = 2.0
 Identities = 47/211 (22%), Positives = 84/211 (39%), Gaps = 32/211 (15%)

Query: 210 IYNLLQKHKNMTSRLYKNIFLEMRNDYVNEKRNTRAFNSNDDAMTTGEFNEYNLADD 269
 I++LLQK LY+N+ + R + N+ T E ++ + ++
 Sbjct: 918 IFSLLQKDSPLLVLYENVHI-----REGEKYGRNE--ATDNEVDYKKGDIKH 964

Query: 270 NLRNHINQNGDFFYIKTD---DKYIKVMYNVTFMTNIIIVPYTKQYEFCTKIRDIDNHV 326
 N+ N + D + D+ K MY + V E K D+ N+
 Sbjct: 965 NVTNEHGNHSDSYFYGNSLNLDRKPKNMYE-DIYKEKGFVKSDCSNIEI--KKNDMINND 1021

Query: 327 TYLRDDMPYKENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKII----KPHIKNE 382
 Y +++ FY+++ Y+ + YV++ +YL +N ++ F +KN+
 Sbjct: 1022 VYKQNE-FYEDSRINMIYDEDEIKTWFLIPHKYVIN---IIYFLNILLTDESFPKLKNK 1077

Query: 383 MKQNMSEFERKEKIYEDN-----YIENTKKY 408
 E K IYEDN ++N KKY
 Sbjct: 1078 KYGYFVNBEETKGTIYEDNNGLEILKNGKKY 1108

Score = 33.6 bits (75), Expect = 2.7
 Identities = 42/198 (21%), Positives = 77/198 (38%), Gaps = 42/198 (21%)

Query: 222 SRLYKNIFLEMR---RNDYVNEKRNTRAF-----NSNDDAMTTGEFNEYNLA 267
 S LY I++ + +N + K+NT + N+++D TT E + +
 Sbjct: 411 SVLYSIYMNKKYKKNFITNKKNTNVYFENDVIQLSVENTSEDITFTTNTRESSLNSGM 470

Query: 268 DDNLRNHINQNGDFFYIKTDCKYIKVMYNVTFMTNIIIVPYTKQYEFCTKIRDIDNHVT 327
 +++R +N D +DDK ++Y N YTK E
 Sbjct: 471 MNDMRYSVNNYADEKVYHSDDKSDHLIYKHVHDEKNKYDEMYTKTKE----- 517

Query: 328 YLRDDMPYKENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKPHIKNEMKKNM 387
 +++ YK N+ + N K LD+ K I H+KN+ + N
 Sbjct: 518 --NENIYKSNIVDKKTCDISSEMVNGKDK-----LDVEKYIGSHVKN-DENN 563

Query: 388 SEFERK-EKIYEDNYIEN 404
 + ++K + + + YI+N
 Sbjct: 564 EKLKKKIDNVNKKKEYIDN 581

>gi|3845297 (AE001421) hypothetical protein [Plasmodium falciparum]
 Length = 2380

Score = 48.0 bits (112), Expect = 1e-04
 Identities = 87/390 (22%), Positives = 160/390 (40%), Gaps = 65/390 (16%)

Query: 20 VRKKIPDKYNQIELVDELMNDDIDYIYSISNRSGKSFNYVSFF-----IYLAIKLDIKF 74
 +++K +K ++ + +N D + ++ R K+ NY++ +YL I DI
 Sbjct: 1049 LQRKNMKNCSKNRNRNRYINKDSNIHLMLIRIKFKNLNMYNMNMFIEIYLLKINNDIFL 1108

Query: 75 TLLSRHYTLRDAYR-----DFIEEIIDEN-PLFKSKRVTPRSARDYLAIYQDKEIGVI 127
 +Y +++ Y + + + EN + +++ ++ + Y +K+
 Sbjct: 1109 QFNKHYNVQNFYNFYSITLINIMSKYSENFYAYNLEKIVYKFLNKNKNFEYIEKQYSSK 1168

Query: 128 TDLNSATDLKYHSNFKHYPIIYDEFLA----LEDDYLIDEWDKLKTIYESIDRNHGNV 183
 D+N D+ ++ +K+ II EFL L+ D I + KLKT ++
 Sbjct: 1169 EDMVEL-DILVNTYDMKYDKII---EFLKNGYLKIDRIYIYFYPKLKT-----DI 1214

Query: 184 DYIGFPMFLGNVNFSSPILSNLNIYNLLQKHKNMTSRLY-----KNIF--LEMRRN 235
 F ++FL N + L NI +++ K + Y K IF + M+ +
 Sbjct: 1215 ILFFPKEIFLNDNILKIDRKFLKK-NITIMIEVLKEIFKEYVKRCITKVIFFPVHMKEH 1273

Query: 236 DYVNEKR-----NTRAFNSNDDAMTTGEFNEYNLADDNLRNHINQNGDFFYIKTD 287
 D+V K N+ FN+ D + N YN D+ N+ N N +Y K

296

Sbjct: 1274 DHVMKNQYNNQYVNSNMFTNRGDHNNNNQTNNDNHNHYYDDTHNNNNNNNSKYK-KNK 1332

Query: 288 DKYIKVMYNTTFTMTNIIV---VPYTKQYEFCTKIRDIDNHVTYLRDDMFYKEN---ME 340
 +K K+MY +++ + V K + K I + Y+ ++ N +

Sbjct: 1333 NKN-KIMYEKERKSSSLFISNNVQDVKPIKHYLYSSIIYKNFYIIEIKNFNNKITKIN 1391

Query: 341 RY-YYNPSNLHFDNAYSKNYVVDNDRYLYL 369
 RY YYN NL+ D+ ND YL+L

Sbjct: 1392 RYNNYNNMNLNIDDL-----NDAYLFL 1413

Score = 32.5 bits (72), Expect = 6.0
 Identities = 46/183 (25%), Positives = 73/183 (39%), Gaps = 26/183 (14%)

Query: 225 YKNIFLEMRRNDYVNEKRNTAFNSNDAMTTGEFEFNEYNLADDNLRNHINQNGDFFYI 284
 +KNI ++ ++N + NSN + + N N+ +N N IN + I

Sbjct: 27 HKNINKNIKNKKFINIDNSNNCNSNSNSNSNNNNNNNNNIVRNN--NNFINADKKKNVI 85

Query: 285 KTDDKYIKVMYNTTFTMTNIIVVPYTKQYEFCTKIRDIDNHVTYLRDDMFYKENMERYYY 344
 +D IK V NI Y ++ + D+ N+ + + KE ER

Sbjct: 86 LNEDDDIKNKELVDESFNIF--YENYFKNLFNLNDVSNKVI--NIIEQKEGDER--- 138

Query: 345 NPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIKFIKNEMKKNMSEFERKEKIYEDNYIEN 404
 N N N +KN V DN +NK IKN +N++E Y N++ +

Sbjct: 139 NADN----NLKNKINVRDN-----INK-----IKN--TRNVNEILIYNNKYIINFLND 180

Query: 405 TKK 407
 T K

Sbjct: 181 TTK 183

>gi|4493936|emb|CAB38972.1| (AL034556) predicted using hexExon;
 MAL3P5.6 (PFC0600w), Hypothetical protein, len: 250 aa
 [Plasmodium falciparum]
 Length = 249

Score = 47.3 bits (110), Expect = 2e-04
 Identities = 53/215 (24%), Positives = 87/215 (39%), Gaps = 30/215 (13%)

Query: 209 NIYNLLQKHKNMNTSRLYKNIFLEMRRNDYVNEKRNTAFNSNDAMTTGEFEF--NEYNL 266
 NIYN L++ YKN N ++ +N N+N EFE N YN

Sbjct: 13 NIYNKLEEK-----YKNFLKLNKNSHMGASQNMNV--NNYTMNELEFEKINNYYNN 64

Query: 267 ADDNLNRNHNQNGDFFYIKTD----DKYIKVMYNTTFTMTNIIVVPYTKQYEFCTKIRD 321
 ++N+ N+IN D+ IK +K ++ YN + I T ++

Sbjct: 65 NNNNINNNINNYDYDMNIKVSQSVQHNRQLQDFYNNKNSFQHYIKKLKTCRFDADDIRNL 124

Query: 322 IDNHVTYLRDDMFYK----ENMERYYYNPSNLHFDNAYSKNYVVDNDRYLYLDMNKIIK 376
 ++ + Y RD+ K EN + N + N+ S NY DN+ LY +N++ K

Sbjct: 125 LEKRLAYERDNTLIKNIQEBENKKGIGINGNFGSESNSSSSNY--DNNYLLYRKINRLNK 182

Query: 377 FHIKNEMKKNMSEFERKEKIYEDNYIENTKKYLMK 411
 + ++ KI KKY++K

Sbjct: 183 TNTNKSKNRSRKRKRINSKI-----DKKYI 209

>gi|3845165 (AE001390) hypothetical protein [Plasmodium falciparum]
 Length = 1247

Score = 45.7 bits (106), Expect = 6e-04
 Identities = 52/239 (21%), Positives = 94/239 (38%), Gaps = 38/239 (15%)

Query: 206 SNLNIYNLLQKHKNMNTSRLYKNIFLEMRRNDYVNEKRNTAFNSNDAMTTGEFEFNEYN 265
 +N N +N ++K K R I +N + +N ++N+D E N N

Sbjct: 474 NNTNKNWBEIKRKKKFKREKNKIINNSFQNEAEDDKNNNNNDNNNDNHNNDNNNNNNEN 533

Query: 266 LADDNLNRNHNQNGDFFYI-KTDDKYIK----VMYNTTFTMTNIIVVPYTKQYEFCTKIR 320
 D+N N+ + N D I D+ Y +YN T ++ YTK + + +

Sbjct: 534 NNDNNNNNNNDINNDINNIHNNNDNNYNNNDNNINLYNEMTKKKCMLDNSYTKYFFYIFTL- 592

Query: 321 DIDNHVTYLRDDMFYKENME-----RYYYN-----PSNLHFDNAYS 356
 + + ++ + FY++N + ++YYN + N

Sbjct: 593 ---DMLPSIKFETFYEKNTDHNKNFNENYKPYNTDDDDTDIINAIKKNVKNKKKNGNIVI 649

Query: 357 KNYVVDNDRYLYLDMNKIIKFIKNEMKKNMSEFER----KEKIYEDNYIENTKKYLMK 411
 KNY+ N+ Y YL+ N+ + I + K +E K+ I+ ++Y E K K

297

Sbjct: 650 KNYINHNE-YSLEYNENKNYEINKKEKLLTENYEYDMYIKDNIHYNDYSEGDGKQTKK 707

Score = 41.0 bits (94), Expect = 0.016
Identities = 58/245 (23%), Positives = 96/245 (38%), Gaps = 43/245 (17%)

Query: 207 NLNIYNLLQKHKMTSRLYKNIFLEMRRNDYVNEKRNTRAFNSNDAMTTGEFEFNEYNL 266
N+N+YN + K K Y F + D + + N D E YN

Sbjct: 564 NINLYNEMTKKKCMLDNSYTKYFFYIFTLDMLPSIKPETFYEKNTDHKNFNENYKFFYNT 623

Query: 267 ADD-----NLRNHINQNGDPF---YIKTDDKYIKVMYNVT-TFMTNIIIVPYTKQ 312
DD N++N +NG+ YI ++ Y + YN + N T+

Sbjct: 624 DDDTDIINAIAKKKNVKNK-KKNGNIVIKNYINHNE-YSLEYNENKNYEINKKEKLLTEN 681

Query: 313 YEFCTKIRDIDNHVYTLRDDMFYKENMERYYYNPSNLHFDNAYS-----NYV--VD 362
YE+ I+D ++ Y D + + YN +N +N Y K +Y+ VD

Sbjct: 682 YEYDMYIKDNIHYNDYSEGDGKQTKKASSFLYNWNN---NNKYKKEDNKTQIISYMDHVD 738

Query: 363 NDR-----YLIDMKNKIKPHIK-NEM---KKNMSEFERKEKIYEDNYIENTKKY 408
N+ Y + ++ F +K N+M K+ F +E I + +EN K+

Sbjct: 739 NENGVKGLKKRNLFFYNNSDQLYNFDVKDNDMIKYEKRQSKNFVEEFINGNRKMNEDKH 798

Query: 409 LMKQY 413
L K Y

Sbjct: 799 LKKHY 803

Query= pt|110877 44AHJDORF007 Phage 44AHJD ORF |2044-3027|1 1
(327 letters)

>gi|1181960|emb|CAA87731.1| (Z47794) connector protein
[Bacteriophage CP-1]
Length = 337

Score = 45.7 bits (106), Expect = 5e-04
Identities = 44/184 (23%), Positives = 84/184 (44%), Gaps = 13/184 (7%)

Query: 127 QIHKLYDNCMSGNFVVMQNKPIQYNSDIEIIEHYTDELAEVALSRFSLIMQAKFSK--IF 184
++HK + + +V+ N Y I +E + ++LA++ L+ L A+ + IF

Sbjct: 125 ELHKDNPDKIKRPCIVIPNNNF-YEPYIGYLELFCEKLADIILT-IQLNRNAQITPYFIF 182

Query: 185 KSEINDESINQLVSEIYNGAPFVKMSPMFNAD-----DDIIDLTSNSVIPALTEMKR 236
N S+ + ++I N P V ++ + D D I + L ++

Sbjct: 183 ADNTNVLSMKNIFNKIANFEPVVVLNKQKQDQDQDSFKQLSDYIQVFRPTDAPFLDLKLHD 242

Query: 237 EYQNKISELSNYLGINSIAVDKESGVSDERAKSNRGFTTSNSNIYLKGREP-ITFLSKRY 295
E +++L ++GIN+ DK+ + EA SN G ++N + K R + ++K Y

Sbjct: 243 EKLVRVNLQLLTFIGINNNPSDKKERLVVSEAISNNGVISANIEVGWKSRRKFVELINKCY 302

Query: 296 GLDI 299
GL+I

Sbjct: 303 GLEI 306

>gi|1429239|emb|CAA67658| (X99260) upper collar protein
[Bacteriophage B103]
Length = 308

Score = 44.9 bits (104), Expect = 8e-04
Identities = 40/159 (25%), Positives = 73/159 (45%), Gaps = 11/159 (6%)

Query: 150 YNSDIEI-----IEHYTDELAEVA-LSRFSLIMQAKFSKIFKSEINDESINQLVSEIYNG 203
YN+D++ +E + +LAE+ + + Q I ++ N S+ + ++

Sbjct: 121 YNNDLKCSSTLPALEMFQDLAELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYQYEGN 180

Query: 204 APFVKMSPMFNADD-DIIDLTSNSVIPALTEMKREYQNKISELSNYLGINSIAVDKESGV 262
AP + + + D+ + + V+ L K N E+ YLGI + ++K+ +

Sbjct: 181 APVIFVHESLDLDNLKVFKTAPYVVDKLNQNAVWV---EVMTYLGIKNANLEKKERM 237

Query: 263 SDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
E SN S+ NIYLK R E +S+ YGL++K

Sbjct: 238 VTSEVDSNDEQIESSGNIYLKARQEACNKISELYGLNLK 276

>gi|137915|sp|P07535|VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR
PROTEIN) (LATE PROTEIN GP10) >gi|75851|pir||WMBP10 gene

10 protein - phage PZA >gi|216059 (M11813). upper collar
protein [Bacteriophage PZA]
Length = 309

Query: 150 YNSDIEI-----IEHYTDELAEVALSRFSFLIMQAKFSKIF--KSEINDESINQLVSEIYN 202
 YN+D+ +E + ELAE+ S+ A+ + + ++ N S+ Q+ ++
 Sbict: 122 YNNDSMFPPTPTTLELFAAEELAEK-EIISVNOQAQKTPVLIRANDNNQLSLKQVYNQYEG 180

Query: 203 GAPFVKMSPMFNADD-DIIDLTNSVIPALTEMKREYQNKISELSNYLGINS LAVDKESG 261
AP + ++D ++ + V+ L K N E+ +LGI + ++K+
Sbjct: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNAQKNAVWN--EMMTFLGIKNANLEKKER 237

Query: 262 VSDEEAKSNRGFTTSNSNIYLKGR-EPITFLSKRYGLDIK 300
+ +E SN S+ ++LK R E +++ YGLD+K
Sbjct: 238 MVTDEVSSNDEOIESSGTVFLKSREEACEKINELYGLDVK 277

```
>gi|137914|sp|P04332|VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR
PROTEIN) (LATE PROTEIN GP10) >gi|75852|pir||WMBPC9 gene
10 protein - phage phi-29 >gi|215328 (M14782) upper
collar protein [Bacteriophage phi-29] >gi|215340
(M12456) p10 connector protein [Bacteriophage phi-29]
>gi|224161|prf||1011232A protein p10.connector
[Bacteriophage phi-29] >gi|225365|prf||13012708 gene 10
[Bacteriophage phi-29]
Length = 309
```

Score = 41.4 bits (95), Expect = 0.009
Identities = 37/160 (23%), Positives = 75/160 (46%), Gaps = 13/160 (8%)

Query: 150 YNSDIEI-----IEHYTDELAEVALSRFSLSIMQAKFSKIF--KSEINDESINQLVSEIYN 202
 YN+D+ +E + ELAE+ S+ A+ + + ++ N S+ Q+ ++
 Sbict: 122 YNNDMAFFPTTPTLELFAAEALAK-EIISVNONAOKTQVLIRANDNNQLSKQVQYNOYEG 180

Query: 203 GAPFVKMSPMFNADD-DIIDLTNSVIPALTEMKREYQNKISELSNYLGINS LAVDKESG 261
 AP + + + + + V + L K N E + + LGI + + + + +
 Sbict: 181 NAPVIFAHEALDSDSIEVFKTDAPYVVDKLNAQINAVNN--EMMTFLGIKIVANLEKKER 237

Query: 262 VSDEEAKSNRGFTTNSNIYLKGR-EPITFLSKRYGLDIK 300
+ +E SN S+ ++LK R E +++ YGL++K
Sbjct: 238 MVTDEVSSNDEQIESGTVFLKSRREEACEKINELYGLNVK 277

Query= pt|110878 44AHJDORF008 Phage 44AHJD ORF |3020-3775|2 1
(251 letters)

```
>gi|4982468|gb|AAD30963.2| (AF118151) SNF1/AMP-activated kinase  
    {Dictyostelium discoideum}  
    Length = 718
```

Score = 52.3 bits (123), Expect = 3e-06
Identities = 28/118 (23%), Positives = 56/118 (46%), Gaps = 5/118 (4%)

[illegible]

Query: 177 TTLRFADNNTIDNGKTVNKSSESQNAKRQNQKGNAGTQFTKQYLID-NIDKAYD 233
 +NN I+N N ++N +N N N N N+ + T+ + I N++ +Y+
 Sbict: 442 NNNNNNNNNIINNNTNNNNNNNNNNNNNNNNNNNNNNSSISGTEVFSISPNNLNNSYN 499

Score = 37.5 bits (85), Expect = 0.094
Identities = 17/111 (15%), Positives = 45/111 (40%)

[illegible]

Query: 190 GKTVKNSSNESQNAKRNQNQGKNAKGTFQTKQYLIDNIDKAYDLRKKILN 240
N +N +N N N N ID+++ + + N
Sbjct: 516 NNNTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNNCIDSVNNSLNNENDVMN 566

Query: 216 GTQFTKQYLIDNIDKAYDLR 235
 Q L++++D D++
 Sbjct: 614 NLNNNFQ-LLNSLDLNSDIQ 632

Query: 186 TIDNGRTVKNSSNESNQAKRNQKGNAGTQTFTKYQLIDNIDKAYDLRKKILN 240
DN N ++N +N N N N N + +++ D+ +N
Sbjct: 520 NNND---NNNNNNNNNNNNNNNNNNNNNNNNNNNNCIDSVNLSLNNENDVNNNSIN 570

Query: 170 VNIDVDNTTLRFADNNTIDNGKT VNKSSNESNQNAKRNQNQKGN 213
 +N ++ + ++ + +N N +++ +N N N N N
 Sbict: 494 LNNSYNSNSSSGNSGNSNNNSNNNTTNDNNNNNNNNNNNNNNNN 537

Query: 190 GKTVNKSSNESNQNAKRNNQKGN 213
+ N +N N N N N
Sbjct: 515 SNNNTNNNNNNNNNNNNNNNNNNNN 538

Query: 186 TIDNGKTVNKSSNESNQNAKRNNQKQNAKGTQFTKQYLIDNIDKA 231
 +N N +SN +N N N N TK+ I + D++
 Sbjct: 502 NNNNNNSNSNSNSNNNNINNNNNNNNNNNIYLTKKPSIGSTDES 547

Query: 87 NRQTV E A P G M Q V I T V C I T H E D Y L N V V Y S S S E V E K Y L Q S Q G F T E H N E D T T S N T D E T S N Q N A 146
N G I T T + + + + + + + N + + N + + N N

300

Sbjct: 415 NNNNNNIIGNGKITTTTTTSTSPSSINNEDISSNNNNNNNNNNNNNNNNNNNNNN 474

Query: 147 TSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESQN 203
+ +++++ T N N + + N + +N N+ +N N ++N +N N

Sbjct: 475 NNNNSNSSNTNNNNINNTTNNNSNSNNNNNNNSNSNSNNNNINNNNNNNNNNN 531

Score = 33.2 bits (74), Expect = 1.8
Identities = 18/88 (20%), Positives = 35/88 (39%)

Query: 130 HNEDTTSNTDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189
+N + ++N + +N N T T + S+ +E +N +NN +N

Sbjct: 405 NNNNSNNNNNNNNNNNIIGNGKITTTTTTSTSPSSINNEDISSNNNNNNNNNNNNNNNN 464

Query: 190 GKTIVNKSSNESQNAKRQKQKGNAGT 217
N ++N +N N+ + N T

Sbjct: 465 NNNNNNNNNNNNSNSSNTNNNNINNT 492

Score = 32.5 bits (72), Expect = 3.1
Identities = 18/94 (19%), Positives = 37/94 (39%)

Query: 120 KYLQSQGFTEHNEDTTSNTDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTL 179
K + S N + +N++ +N N ++ + +T S N D+ +

Sbjct: 392 KNVVSTSLVPGNGNNNNNNNNNNNNNNNNNNNIIGNGKITTTTTTSTSPSSINNEDISSNNN 451

Query: 180 RFADNNTIDNGKTVNKSSNESQNAKRQKQKGN 213
+NN +N N ++N +N N + + N

Sbjct: 452 NNNNNNNNNNNNNNNNNNNNNNNNNNSNSSNTN 485

Score = 32.5 bits (72), Expect = 3.1
Identities = 24/110 (21%), Positives = 44/110 (39%), Gaps = 10/110 (9%)

Query: 138 TDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGK----- 191
T T++ + +S++N+ ++N N + + N + +N +NN N

Sbjct: 429 TTTTSTSPSSINNEDISSNNNNNNNNNNNNNNNNNNNNNNNNNSNSSNTNNNN 488

Query: 192 ----TVNKSSNESQNAKRQKQKGNAGTQFTKQYLIDNIDKAYDLRKK 237
T N +SN +N N N N N+ +N + L KK

Sbjct: 489 INNTTNNNSNSNNNNNNNSNSNSNNNNINNNNNNNNNNNIYLTKK 538

>gi|3758855|emb|CAB11140.1| (Z98551) predicted using hexExon;
MAL3P6.11 (PFC0760c), Hypothetical protein, len: 3395 aa
[Plasmodium falciparum]
Length = 3394

Score = 46.5 bits (108), Expect = 2e-04
Identities = 52/202 (25%), Positives = 96/202 (46%), Gaps = 32/202 (15%)

Query: 21 FNEFVNDNKLTIFYDDEFQFMQKMLKFD-KDVLAIVNEKVFKGFSKDELSL--LFKKSF 77
F ++ ++ K T D+ M+K K D DV + NEK++ L ++L+ + + KK

Sbjct: 665 FEKYCSNIKNTLIRDD---MKKFRKPDISDVHILHNEKIYLEKLLNEKLNYYIKDIEKKLD 721

Query: 78 TIHFLDREINRQTVEAFGMQV-----ITVCITHEDYLNVVYSSEVEKYLSQGFTEHNE 132
+H + IN+ + + +QV I V + DY + S + + K + +N

Sbjct: 722 ELHGV---INKNKEDIYILQVEKQTLIKVISSVYDYTKME-SENHIFKMTTWNKMLNNV 777

Query: 133 DTTSTDETSNQATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKT 192
+SN D +NQN ++N+ + N+N N +++N + N +N

Sbjct: 778 HSSNKDY-NNQNNQNIENNQNIENNQN-----NQNIEN-----NQNIENNQN 820

Query: 193 VNKSSNESQNAKRQKQKGN 214
N +N++NQN + NQN + NA

Sbjct: 821 QNNQNNQNNQNNQNNQNNQNNNA 842

Score = 33.6 bits (75), Expect = 1.4
Identities = 46/221 (20%), Positives = 89/221 (39%), Gaps = 37/221 (16%)

Query: 10 DFIKSELIKKGFNEFVNDNKLTIFYDDEFQFMQKMLKFDKDVLAIVNEKVFKGFSKDELS 69
D +K E K N + +L Y + + M+K K + V K SL

Sbjct: 367 DSLKIEYNKSKTNIQQLNEQLVNYKNFIKEMEKKYK-----QLVVKNNSLFSITH 416

Query: 70 DLLFKKSFTIHFLLDREINRQTVEAFGMQVITVCITH---EDYLNVVYSSEVEKYLSQSG 126

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      D + K+ I + R + + +      ++ + I H      +D+L+V+Y      + + L +
Sbjct: 417 DFINKLNSNIIIRRTSDMKQI----FKMYNLDIEHFNEQDHLSVIY----IYEILYNTN 468

Query: 127 FTEHNEDTTSNTDETSNQNAATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
      +N D +N D +N N + +N+      N N      N + +N +
Sbjct: 469 -DNNNNDNDNNNDNNNNNNNNNDNNNNNNNDNNNN-----NNNYNNIMM-----M 512

Query: 187 IDNGKTVKNSSNESNQAKRNQWQKGNAGTQFTKQYLLDN 227
      I+N + N ++ + + N + N + N + + + Y I+N
Sbjct: 513 IENMNSGNHPNSNNLHNYRHNTDNENNLSLKTFSFRYKINN 553

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Query: 119  EKYLQSQGTFEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID-VDNT 177
           E Y S + +++ N + +N + + DN+ N N ++ +N D ++N
Sbjct: 2838  ENYPVSTHYDNDDINKDNINNDNNNDNINDDNNNDNINNDNNNDNINDDNINNDNINND 2897

Query: 178  TLRFADNNTIDNGKTVNKSSNESNQNAKRNNQKGNAGTQFTKQYLIDNIDKAYDLRKK 237
           +N+ +NG SSN ++ N N N K N +G + + + + YD K
Sbjct: 2898  NNNDNNNDNSNNGFVCELSSNINDFNNILNVN-KDNFQGINKSNNFSTNLSEYNIDYAVK 2956

Query: 238  IL 239
           I+
Sbjct: 2957  IV 2958

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Query: 9      YDFIKSELIKGFNEFVNDNKLTPYDDEQFQMQLKPKDKDLAIVNEKVFGKFSLKDEL 68
             Y+++K  ++  N  N  NK      E  Q++ K+  +  +  +E  K  L++
Sbjct: 2150  YNYVK---VQNATNREDNKNK-----ERNLSQEITYKYINENIDLTSELEKQNDMLenYK 2200

Query: 69     SDL-----LFFKSFTIHFLDREINRQTVEAFGMQVITVCITHEDYLVNVVYSSSEVEKYL 122
             ++L  ++K +  I  L      +  M+  +  +      N  +  E+  +  L
Sbjct: 2201  NELKEKNEEYIKLNNDDMLSNCKKLKESIMMEKYKIIMN----NNIQEKDEIIEINL 2255

Query: 123    QSQGFTEHNEDDTTSNTDETSQNQATSLDNSTGMTAN----RNAYVSLPQSE----VNIDV 174
             +++ +      +D  +N      +  ++S  M+  +      N  +  +L  +S      N+D+
Sbjct: 2256  KNK-YNNKLLDDLNNYSVVDKSIYSCFEDSNIMSPSCNDILNVFNNLKSKNKKVCTNMMDI 2314

Query: 175    DNTTLRFADNNTIDNGKTVNKSSNESQNAKRNQNKQNAKGTQFTKQYLIIDNIDKAYDL 234
             N  +      ++I+N  +N  +N  +N  N  N  N  K      YL++N+  D
Sbjct: 2315  CNEIMDSI--SSINNVNNINNVNNINNVNNINNVNNINNVKNIVDINNYLVNNLQLNKDN 2372

Query: 235    RKKILNEFD 243
             I+  +F+
Sbjct: 2373  DNIIIIKFN 2381

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Query: 115 SSEVEKYLQSQGFTEHNEDDTTSNTDETSQN--ATSLDNSTGMTANRNAYVSLPQSEVNI 172
+++ EK Y EH + N D +N+N L ++ ++ + N S ++E+
Sbjct: 3264 NNDEEKYSCHDDKNEHTNNDLLNIDHDNNKNNITDELSTYNVSVSHNKDPSNKEINQ 3323

Query: 173 DVDNTTLRFADNNITIDNGKTVNKSSESNEQNAKRNQNKGNK 215
+ + D N ++ N ++E+++N + ++N + + K
Sbjct: 3324 LLSIDSSNENDENDENDENDENDENDENDENDEK 3366

Query: 104 THEDYLNVVYSSEV----EKYLSQSGFTEHNEDTTSNTDETSQNQATSLDNSTGMTANR 159
T+ D LN+ + +++ E Y HN+D ++ +E QN S+D+S N
Sbjct: 3280 TNNDLLNIDHDNNKNNITDELYSTYNVSVSHNKOPSNKENEI--QNLISIDSSNENDEND 3337

Query: 160 NAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVKNKSSNESNQNAFRNQNKQKNAKT 217
+++ N + D D N ++ N +E+++N + ++ N +GT
Sbjct: 3338 EN----DENDENDEND-----DENDENDENDEKDEKNDENDENDENFDNNNEG 3386

302

>gi|585795|sp|P21538|REB1_YEAST DNA-BINDING PROTEIN.REB1 (QBP)
 >gi|626139|pir|S45907 DNA-binding protein REB1 - yeast
 (Saccharomyces cerevisiae) >gi|536280|emb|CAA84992|
 (235918) ORF YBR049c [Saccharomyces cerevisiae]
 >gi|559944|emb|CAA86391| (Z46260) REB1 DNA-binding
 protein [Saccharomyces cerevisiae]
 Length = 810

Score = 45.7 bits (106), Expect = 3e-04
 Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)

Query: 83 DREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETS 142
 D+ N+++VE ++ + V + H+++ +++ K+ + Q E + D N ++ S
 Sbjct: 7 DKNANQESVEEAVLKYGVLGHQNHDPQLHTKDLENKHSKKQNIVESSSDQVDVNNNDSS 66

Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
 N+N + D+S ++A L +E + +VD+ N +D N+ +E
 Sbjct: 67 NRNEDNNDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119

Query: 200 SNQNAKRNNQKGNAGTQFTKQYLIDNIDKAYDLRKK 237
 ++N N GN F++ ++ +D D KK
 Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153

>gi|172372 (M58728) DNA-binding protein [Saccharomyces cerevisiae]
 Length = 809

Score = 45.7 bits (106), Expect = 3e-04
 Identities = 34/158 (21%), Positives = 72/158 (45%), Gaps = 14/158 (8%)

Query: 83 DREINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLQSQGFTEHNEDTTSNTDETS 142
 D+ N+++VE ++ + V + H+++ +++ K+ + Q E + D N ++ S
 Sbjct: 7 DKNANQESVEEAVLKYGVLGHQNHDPQLHTKDLENKHSKKQNIVESSSDQVDVNNNDSS 66

Query: 143 NQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID---NGKTVNKSSNE 199
 N+N + D+S ++A L +E + +VD+ N +D N+ +E
 Sbjct: 67 NRNEDNNDSENISA-----LNANESSSNVDHANSNEQHNAVMDWYLRQTAHNQQDDE 119

Query: 200 SNQNAKRNNQKGNAGTQFTKQYLIDNIDKAYDLRKK 237
 ++N N GN F++ ++ +D D KK
 Sbjct: 120 DDEN--NNNTDNGNDSNNHFSQSDIV--VDDDDDKNKK 153

>gi|2952545 (AF051898) coronin binding protein [Dictyostelium
 discoideum]
 Length = 560

Score = 44.9 bits (104), Expect = 6e-04
 Identities = 26/83 (31%), Positives = 39/83 (46%), Gaps = 5/83 (6%)

Query: 131 NEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNG 190
 N + +N +N N+ S +NS +N N+ + P N D DN T +NNT +N
 Sbjct: 404 NNNNNNNIINNNSNSNSNNNSNN-NSNNNSNRNSPNHNNNGDNDNNT----NNNTNNNN 458

Query: 191 KTVNKSSNESQNAKRNNQKGN 213
 N ++N +N N N N N
 Sbjct: 459 NNNNNNNNNNNNNNNNNNNNNNNNNNNNN 481

Score = 41.4 bits (95), Expect = 0.006
 Identities = 22/88 (25%), Positives = 43/88 (48%), Gaps = 6/88 (6%)

Query: 130 HNEDTTSNTDETSNQNATSLDN---STGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNT 186
 + ++ +N++ SN N+ + +N + G AN++ + P + +N + DN +NN
 Sbjct: 337 NRNNSNNNSNNNSNNNSNNNRNITNGSNANKS---NSPNNNLNTNNDNKNNSNNNNNN 393

Query: 187 IDNGKTVNKSSNESQNAKRNNQKGN 214
 +N S+N +N N N N N+
 Sbjct: 394 SNNNSNNGNSNNNNNNNNIINNNSNSNS 421

Score = 40.6 bits (93), Expect = 0.011
 Identities = 24/101 (23%), Positives = 41/101 (39%), Gaps = 2/101 (1%)

Query: 115 SSEVEKYLQSQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDV 174
 S+ L + ++N +N ++ N S +N+ N N S + N +

Sbjct: 370 S N S P N N N L N T N N D N K N N N S N N N N N S N N N S N N G N S N N N N N N N I I N N N N S N S N S N N N S N N N S 429

Query: 175 DNTTLRFADN--NTIDNGKTVNKSSNESNQNAKRNQNQKGN 213
+N + R + N N DN N ++N +N N N N N

Sbjct: 430 NNNSNRNSPNHNNNGDNDNNTNNNTNNNNNNNNNNNNNNNNNN 470

Score = 40.2 bits (92), Expect = 0.014
Identities = 21/80 (26%), Positives = 39/80 (48%), Gaps = 9/80 (11%)

Query: 130 HNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDN 189

Sbjct: 442 +N D +NT+ +N N + +N+ N N N + +N +ADN+ ++
NGGDNDNNNTNNNTNNNNNNNNNNNNNNNNNN-----NNNNNNNNNYADNSNNNS 492

Query: 190 GKTVNKSSNESNQNAKRNQN 209

Sbjct: 493 SNSNNNSNSNNNDNKNEN 512

Score = 39.5 bits (90), Expect = 0.024
Identities = 26/111 (23%), Positives = 44/111 (39%), Gaps = 20/111 (18%)

Query: 112 VYSSSEVEKYLQSQ--GFTEHNEDTTSNTDETSNQATSLDNSTGMTANRNAYVSLPQSE 169
 VY + K+ ++ G +N ++ +N++ SN N ++N N N

Sbjct: 296 VYCTHHHTKFYETHRGLNNNNNSNNNSNSNNNNNGINNRRNSNNNSN----- 346

Query: 170 VNIDVDNTTLRFADNNTIDNGKTVNKSS-----NESNQNAKRNQNQKGNA 214

Sbjct: 347 ---NNSNNNSNNSNNRNITNGSNANKSNSPNNNLNTNNDNKNNNSNNNNNS 394

Score = 37.5 bits (85), Expect = 0.094
Identities = 24/96 (25%), Positives = 41/96 (42%), Gaps = 1/96 (1%)

Query: 124 SQGFTEHNEDTTSNTDETSNQATSLDNSTGM-TANRNAYVSLPQSEVNIDVDNTTLRFA 182
S + +N + SN + ++ N DN+T T N N + + N + +N

Sbjct: 421 SNNNSNNSNNSNRNSPNHNNNGDNDNTNNNTNNNNNNNNNNNNNNNNNNNNNNNNNNNNNN 480

Query: 183 DNNTIDNGKTVNKSSNESNQNAKRNQNQKGNAKGTQ 218

Sbjct: 481 NNNYADNSNNNSSNSNNNSNSNNNDNKNENSDNQ 516

Score = 35.6 bits (80), Expect = 0.36
Identities = 25/99 (25%), Positives = 42/99 (42%), Gaps = 18/99 (18%)

Query: 130 HNEDTTSNTDETSNQNATSLDNST-GMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTID 188

Sbjct: 339 NNSNNNSNNNSNNNSNNNSNNRNITGNSNANKS--NSPNNNLNTNNDNKNNNSNNNNNSN 395

Query: 189 NGKTV-----NKSSNESNQNAKRNQNQKGN 213

N N S++ SN N+ N N N
Sbjct: 396 NNSNNGNSNNNNNNNNIINNNSNSNSNNNSNNNSNNNSN 434

Score = 35.2 bits (79), Expect = 0.47
Identities = 21/94 (22%), Positives = 42/94 (44%), Gaps = 5/94 (5%)

Query: 124 SQGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNIIDVNTTLRFAD 183
+ G + ++ +N T+N N + N+ N N+ + N + +N + +

Sbjct: 362 TNGSNANKSNSPNNLNTNNDNKNNNSNN-----NNNSNNNSNNGNSNNNNNNNNIINN 416

Query: 184 NNTIDNGKTVNKSSNESNQNAKRNQNQKGNAGT 217

+N+ N + N S+N SN+N+ + N N T
 Sbjct: 417 SNSNSNNNSNNNSNNNSNRNSPNHNNNGDNDNNT 450

Score = 35.2 bits (79), Expect = 0.47
Identities = 29/118 (24%), Positives = 53/118 (44%), Gaps = 12/118 (10%)

Query: 115 SSEVEKYLS-QGFTEHNEDTTSNTDETSNQNATSLDNSTGMTANRNAYVSLPQSEVNID 173

SS+ E ++ +GF + + T+N ++N D S+G + + + V+ P+S +N
 Sbjct: 114 SSDSEADIEDDKGFQD--KPITTNNSGSNNPLKLNLDYSSGSSGSSRSQVNPQRSNINNS 171

Query: 174 VDN⁺TLRFADNNT-----IDNGKTVNKSSNESNQNAKRNQ⁺NQKGNAGTQFTKQ 222

Sbjct: 172 NDKYKSKSSSSNSNSSSSGGSLISSLLTGGNTYQNQNQNQNQNQNQNQNNNSQLQQQQQ 229

Query: 131 NEDTTSTNDETSNQATSLDSTGMTANKRAYVSLPQSEVNIDVDNTTLRFADNNNTIDNG 190
N + T + N + + N N + + + N N S N N + N + N
Sbjct: 451 NNNNTNN 504
NN

Query: 124 SQGFTEHNEEDTTSNTDSTSNQNAATSLDNSTGMTANRNAYVSLPQSEVNI DVNTTLRFAD 183
S N SN +++++ N N+ N N + + N + +N
Sbjct: 353 SMNSNNRRNITNGSNANKSNSPNNNLNTNNDKNKNNNSNNNNNSNNNSNNGNSNNNNNNNNII 412

Query: 128 TEHNEDTTSNTDETSNQNATSLDNSTGMTANRRAYVSLPQSEVN-----IDVDNTTLRF 181
T+nnn T TD + + +nn+T A Nn + ++ N D +NT +

Sbjct: 433 TDNNNTNTKATDSNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 492

Query: 128 TEHNEDTTSNTDETSNQNATSLDSS----TGMTANRNAYVSLPQSEVN----IDVDNTTL 179
T++N T TD++N+ + DN+ T T N N S D +NT

Sbjct: 401 TDNNNTDTKATDKSNNTDTKATDNNNNNTDKATDNNNNNTKATDSNNNTKATDNNNTNT 460

Query: 128 TEHNEDTTSNTDETSQNQATSLD-NSTGMTANRNAYVSLPQSEVNIIDVNTNTLRFADNNNT 186
T++N++T++ ++N+N++D+N+T A D N++ ++N NT++DNN

Sbjct: 422 TDNNNNNTDKATDNNNNNTKATDSNNNTKADNNNNNTKATDNN---NTNTKATDNNN 477

Query: 85 EINRQTVEAFGMQVITVCITHEDYLNVVYSSSEVEKYLSQSGFTEHNEEDTTSNTDETSQ 144
E N + + G T + + N + E + + Q T + N T T + + N
Sbjct: 118 ETNKNIKLGTGNNSTINTNLNTENTNA--TKKLTENVNTQILGTGNNNTTNTTSSTEHN 175

Query: 145 NATSLDNSTGMTANRNAYVSLPQSEVNIDVDNTTLRFADNNTIDNGKTVNKSSNESNQNA 204
N + NSTG T+ NI + N L +N T + T + ++ +N N+

305

Sbjct: 176 NINTNTNSTGNTSTTKKLT-----NI-ITNQILTGNNTTTNTSSTEHNNNINTNTNS 228

Query: 205 KRQKQKGNAGTQFTKQYLIDNIDKAYDL 234
 N N N T + DNI+ +L

Sbjct: 229 TDNSNTNTNLTDITTTTKKWDNINTTQNL 258

Score = 41.8 bits (96), Expect = 0.005
 Identities = 30/101 (29%), Positives = 43/101 (41%), Gaps = 13/101 (12%)

Query: 130 HNEDTTSNTDETSNQATSLDNTGTMANRNAYVSLPQSEVNIDV-----DNTTLRFA 182
 +N DT S ++ ++ AT DN+ T T N N + N D +NT +
 Sbjct: 363 NNTDTISTDNDNTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNNTDTKAT 422

Query: 183 DNN-----TIDNGKTVNKSSNESQNAKRQKGNAGKT 217
 DNN DN T K+++ +N N K N N K T
 Sbjct: 423 DNNNNTDTKATDNNNTNTKATDSNNNTNTKATDNNNTNTKAT 463

Score = 40.6 bits (93), Expect = 0.011
 Identities = 31/121 (25%), Positives = 47/121 (38%), Gaps = 31/121 (25%)

Query: 128 TEHNEDTTSNTDETSNQAT-----SLDNTGTMANRNAYVSLPQSEVN----- 171
 TEHN + +NT+ T N + T ++ + +T N N + +E N
 Sbjct: 171 TEHNNNINTNTNSTGNTSTTKKLTENIITNQILTGNNTTTNTSSTEHNNNINTNTNSTD 230

Query: 172 -----IDVDNTTLRFADN-----NTIDNGKTVNKSSNESQNAKRQKGNAGK 216
 D+ TT ++ DN T N TV+ +N +N N K N N K
 Sbjct: 231 NSNTNTNLTDITTTTKKWDNINTTQNLTTSTNTTTVSTDNNNNINTKPTDNNNTNIKS 290

Query: 217 T 217
 T
 Sbjct: 291 T 291

Score = 38.3 bits (87), Expect = 0.055
 Identities = 28/98 (28%), Positives = 41/98 (41%), Gaps = 10/98 (10%)

Query: 128 TEHNEDTTSNTDETSNQATSLDNTGTMANRNAYVSLPQSEVNIDVD-NTTLRFADNNT 186
 TEHN + +NT+ S N+ + N T +T + + N+ NTT DNN
 Sbjct: 216 TEHNNNINTNTN--STDNSNTNTNLTDITTTTKKWDNINTTQNLTTSTNTTTVSTDNNN 273

Query: 187 -----IDNGKTVNKSSNESQNAKRQKGNAGKT 217
 DN T KS++ N K N+ + K T
 Sbjct: 274 NNINTKPTDNNNTNIKSTDNYNTGKETDNKNTDIKAT 311

Score = 37.5 bits (85), Expect = 0.094
 Identities = 31/106 (29%), Positives = 45/106 (42%), Gaps = 18/106 (16%)

Query: 128 TEHNEDTTSNTDETSNQAT-----ATSLDNTGTMANRNAYVSLPQSEVN-----IDVDN 176
 T++N +T +T T N N AT N+T A N + ++ N D +N
 Sbjct: 390 TDNNMT--DTKATDNNNTDTKATDKSNNNTDTKATDNNNTDTKATDNNNTNTKATDSNN 447

Query: 177 TTLRFADNN-----TIDNGKTVNKSSNESQNAKRQKGNAGKT 217
 T + DNN DN T K+++ +N N K N N K T
 Sbjct: 448 TNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKAT 493

Score = 35.2 bits (79), Expect = 0.47
 Identities = 24/109 (22%), Positives = 46/109 (42%), Gaps = 6/109 (5%)

Query: 128 TEHNEDTTSNTDETSNQATSLDNTGTMANRNAYVSLPQSEVN-----IDVDNTTLRF 181
 T++N T TD + + +N+T A N + ++ N D +NT +
 Sbjct: 473 TDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKATDNNNTNTKA 532

Query: 182 ADNNITIDNGKTVNKSSNESQNAKRQKGNAGTQFTKQYLIDNIDK 230
 DNN N + +E+ + K N++ N++ + K + +DK
 Sbjct: 533 TDNNNTNQYVFANNYDETTSDDKLNKDCDENSEEKENIKSMINAYLDK 581

Score = 34.4 bits (77), Expect = 0.81
 Identities = 26/126 (20%), Positives = 46/126 (35%), Gaps = 7/126 (5%)

306

Query: 99 ITVCITHEDYLVVYSSSEVEKYLSQSGFTEHNEDTTSNTDETSNQATSLDNSTGMTAN 158
 IT T+ + ++ S + V S T +++ +N T N N ++ T
 Sbjct: 318 ITTDNTNTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVISTDNNNTDTISTDNDNTDT 377

Query: 159 RNAYVSLPQSEVNIDVDNTTLRFADNNTID-----NGKTVNKSSNESNQAKRNQK 211
 + ++ + +NT + DNN D N + N +N + K N
 Sbjct: 378 KATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKATDNN 437

Query: 212 GNAKGT 217
 N K T
 Sbjct: 438 TNTKAT 443

Score = 34.4 bits (77), Expect = 0.81
 Identities = 30/100 (30%), Positives = 44/100 (44%), Gaps = 14/100 (14%)

Query: 131 NEDTTSNTDETSNQATSLDNTS-TGMTANRNAY---VSLPQSEVNI---DVDNTTLRFAD 183
 N + T TD T N N S DNS T + + N+ +S S+ N+ D +NT D
 Sbjct: 313 NNNITITDNT-NTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVISTDNNNTDTISTD 371

Query: 184 NNTIDNGKTVNKSS-----NESNQAKRNQKGNKGT 217
 N+ D T N ++ N +N + K N + K T
 Sbjct: 372 NDNTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKAT 411

Score = 34.4 bits (77), Expect = 0.81
 Identities = 28/101 (27%), Positives = 41/101 (39%), Gaps = 15/101 (14%)

Query: 131 NEDTTSNTDETSNQATSLDNTS-TGMTANRNAYVSLPQSEVNIDV-----DNTTLRFA 182
 N DT + ++ ++ AT +N+T A N N N D +NT +
 Sbjct: 374 NTDTKATDNDNTDTKATDNNNTDTKATDNNNTDTKATDKSNNTDTKATDNNNTDTKAT 433

Query: 183 DNNITIDNGK-----TVNKSSNESNQAKRNQKGNKGT 217
 DNN N K T K+++ +N N K N N K T
 Sbjct: 434 DNNN-TNTKATDSNNNTKATDNNNTKATDNNNTKAT 473

Score = 32.5 bits (72), Expect = 3.1
 Identities = 30/110 (27%), Positives = 40/110 (36%), Gaps = 23/110 (20%)

Query: 131 NEDTTSNTDETSNQATSLDNTS-TGMTANRNAYVSLPQS-----EVNIDVDNTTLRF 181
 N +TT N ++N S DN+ T T N N + + D NT ++
 Sbjct: 251 NINTTQNLTTSTNTTIVSTDDNNNNINTKPTDNNNTNIKSTDNNTGTKETDNKNTDIKA 310

Query: 182 ADNNTI-----DNGKTVNKSSNESNQAKRNQKGNKGT 217
 DNN I DN KT S + SN + N K N T
 Sbjct: 311 TDNNNITITDNTNTNVISTDNSKTNVISKDNSNTHTISTDNSKTNVIST 360

>gi|1429240|emb|CAA67659| (X99260) lower collar protein
 [Bacteriophage B103]
 Length = 293

Score = 43.8 bits (101), Expect = 0.001
 Identities = 53/204 (25%), Positives = 79/204 (37%), Gaps = 42/204 (20%)

Query: 56 EKVFKEG----FSLKDELSDLLFKKSFTIHFLD---REINRQTVEAFGMQVITVCITHED 107
 EK+ KG F + + D ++K F HF+ REI +T F + T I +
 Sbjct: 26 EKIEKGRPKLFDQYPIFDESRYKVFETHFIRNFYMRIGFETEGLFKFNLETWLIINMP 85

Query: 108 YLVVYSSSEVEKY-----LQSQGFTEH-----NEDTT-----SNTDETSNQNA 146
 Y N ++ S E+ KY L + G ++ N DTT SNT + NA
 Sbjct: 86 YFNKLFES-ELIKYDPLENTRLNTTGNKKNDTERDNRDTTGSMDKGSNTKTSKDTNA 144

Query: 147 TSLDNSTGMTA-----NRNAYVSLPQSEVNIDVDN--TLRFADNNTIDNGKTVNKS 196
 T G T NR P S +N+ ++ TL +A + I+ T NK
 Sbjct: 145 TGSSKEDGKTTGSDVDNFRKIDSDQPSRLNLTNDGQGTLEYA--SAIEENNTNNKR 202

Query: 197 SNESNQAKRNQKGNKGTQFT 220
 + N + + GT T
 Sbjct: 203 NTTGTNNVTSSAESESTGSGTSDT 226

Query= pt|110879 44AHJDORF009 Page 44AHJD ORF |5744-6496|2 1
 (250 letters)

307

>gi|2764981|emb|CAA69021.1| (Y07739) N-acetylmuramoyl-L-alanine
amidase [Staphylococcus phage Twort]
Length = 467

Score = 180 bits (452), Expect = 1e-44
Identities = 89/157 (56%), Positives = 109/157 (68%), Gaps = 8/157 (5%)

Query: 1 MKSQQQAKEWIKHEGAGVDFDGYGFCMDLSVAYVYITDGKVRMNGNAKDAINNDFK 60
MK+ +QA+ +I G DFDG YG+QCMDL+V Y+Y++TDGK+RMWGNKDAINN F
Sbjct: 1 MKTLKQAESYIKSVKNTGTDFDGLYGYQCMDLAVDYIYHVTGKIRMGNAKDAINNNSFG 60

Query: 61 GLATVYKNTPSFKPQLGDVAVYTTNGQ---YGHICVLS----GNLDYYTCLEQNLGGGF 113
G ATVYKN P+P+P+ GDV V+T G YGHI V + G+L Y T LEQNW G G
Sbjct: 61 GTATVYKNYPAPFRPKYGDVVVWTTGNFATYGHIAIVTNPDPYQDLQYVTVLEQNWNGNGI 120

Query: 114 DGWEKATIRTHYYDGVTHFIRPKFSGSNS-KALETSK 149
E ATIRTH Y G+THFIRP F+ +S K +T K
Sbjct: 121 YKTELATIRTHDYTGITHFIRPNFATESSVKKKDTKK 157

Score = 61.7 bits (147), Expect = 6e-09
Identities = 41/125 (32%), Positives = 57/125 (44%), Gaps = 8/125 (6%)

Query: 125 YYDGVTHFIRPKFSGSNSKALETSKVNTEFGKWRNQGTYTYRNENGTFTC-GFLPIFARV 183
YY+G T P +K + +T G W N YGTY+ +E+ TF C I R
Sbjct: 346 YYEGKTPV--PTVVNQAKTKPKVQSSTSG-WNVNNGTYTKSESATFKCTARQGIVTRY 402

Query: 184 GSPKLSEPNQYWFQPNQYTPYNEVCLSDGVVWIGYNWQGR-YYLPVRQWNGKTGNSYSV 242
P + P Y+ VC DGYVWI + G + ++PVR W+ N+ +
Sbjct: 403 TGPFTTCQAGVLYYGQSVTYDTVCKQDGYVWISWTNGGQDVWMPVRTWD---KNTDIM 459

Query: 243 GIPWG 247
G WG
Sbjct: 460 GQLWG 464

>gi|113675|sp|P24556|ALYS_STAAU AUTOLYSIN
(N-ACETYLMURAMOYL-L-ALANINE AMIDASE)
>gi|79887|pir||JQ1147 N-acetylmuramoyl-L-alanine amidase
(EC 3.5.1.28) - Staphylococcus aureus >gi|153067
(M76714) peptidoglycan hydrolase [Staphylococcus aureus]
Length = 481

Score = 118 bits (292), Expect = 6e-26
Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)

Query: 135 PKFSGSNSKALETSKVNTEFGK-WKRNQYGTYYRNENGTFTCGFLPIFARVGSPLSEPNQ 193
P + SN + ++ V WKRN+YGTYY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPIITVRKVGPFPLSCPVG 424

Query: 194 YWFQPNQYTPYNEVCLSDGVVWIGYNWQGRYYLPVRQWNGKTGNSYSVGPWGVFS 250
Y FQ P GY Y EV L DG+VW+GY W+G RYYP+R WNG + +G WG S
Sbjct: 425 YQFPFGGYCDYTEVMLQDGHVWVGTYWEGQRYLPPIRTWNGSAPPNQILGDLWGEIS 481

Score = 78.0 bits (189), Expect = 7e-14
Identities = 48/109 (44%), Positives = 62/109 (56%), Gaps = 6/109 (5%)

Query: 15 EGAGVDFDGYGFCMDLSVAYVYITDGKVRMNGNAKDA-INNDFKGLATVYKNTPSFK 73
EG + D YGFQC D + A + + G + AKD N+F GLATVY+NTP F
Sbjct: 18 EGKQFNVDLWYGFQCFDYANAG-WKVLFGLLKGLGAKDIPFANNFDGLATVYQNTPDFL 76

Query: 74 PQLGDVAVYTTNGQ---YGHICVLSGNLDYYTCLEQNLGGGF-DGWEK 118
Q GD+ V+ + YGH+ V+ LDY EQNLWGGG+ DG E+
Sbjct: 77 AQPQDMVVFSGSYGAGYGHVAVVIEATLDYIIVYEQNLWGGGWDGIEQ 125

>gi|1763243 (U72397) amidase [bacteriophage 80 alpha]
Length = 481

Score = 118 bits (292), Expect = 6e-26
Identities = 56/117 (47%), Positives = 68/117 (57%), Gaps = 1/117 (0%)

Query: 135 PKFSGSNSKALETSKVNTEFGK-WKRNQYGTYYRNENGTFTCGFLPIFARVGSPLSEPNQ 193
P + SN + ++ V WKRN+YGTYY E+ FT G PI R P LS P G
Sbjct: 365 PVATVSNESASSNTVKPVASAWKRNKYGTYYMEESARFTNGNQPIITVRKVGPFPLSCPVG 424

Query: 194 YWFQPNGYTFYNEVCLSDGYVWIGYNWQGTRYLPVRQWNGKTGNSYSGIPWGVFS 250
 Y FQP GY Y EV L DG+VW+GY W+G RYYP+R WNG + +G WG S
 Sbjct: 425 YQFPQGGYCDYTEVMLQDGHVWVGVTWEGQRYLPVRTWNGSAPPNQILGDLWGEIS 481

Score = 83.5 bits (203), Expect = 2e-15
 Identities = 50/115 (43%), Positives = 65/115 (56%), Gaps = 6/115 (5%)

Query: 9 EWIYKHEGAGVDFDGYGFCMDLSVAYVYYITDGKVRMWGNAKDA-INNDFKGLATVYK 67
 EW+ EG + D YGFQC D + A + + G + AKD N+F GLATVY+
 Sbjct: 12 EWLKTSEKQFNVDLWYGFQCFDYANAG-WKVLFGLLKGLGAKDIPFANNFDGLATVYQ 70

Query: 68 NTPSFKPQLGDAVYVYTNQ---YGHQCVLSGNLDYYTCLEQNWLGSGF-DGWEK 118
 NTP F Q GD+ V+ + YGH+ V+ LDY EQNWLGSG+ DG E+
 Sbjct: 71 NTPDLAQPDMVVFSGSYGAGYGHVAVVIEATLDYIIVYEQNWLGSGWTDGIEQ 125

>gi|4574237|gb|AAD23962.1|AF106851_1 (AF106851) *LytN* [*Staphylococcus aureus*]
 Length = 383

Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)

Query: 15 EGAGVDFDGYGFCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
 E G DFDG+YG+QC DL Y ++ ++ +G N+F A +Y NTP+FK
 Sbjct: 252 ENRGWDFDGSYGWQCFDLNVVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311

Query: 75 QLGDVAVYT---NGQYGHQCVLSGNLD---YYTCLEQNWLGSGFDGWEKATIRTHYYD 127
 + GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
 Sbjct: 312 EPGDLVVFSGRFGGGYGHTAIVLNGDYDGKLMKFQSLDQNWNNGGWRKAEVAHKVHNYE 371

Query: 128 GVTHFIRP 135
 FIRP
 Sbjct: 372 NDMIFIRP 379

>gi|3767593|dbj|BAA33856.1| (AB015195) *LytN* [*Staphylococcus aureus*]
 Length = 383

Score = 84.3 bits (205), Expect = 9e-16
 Identities = 48/128 (37%), Positives = 68/128 (52%), Gaps = 7/128 (5%)

Query: 15 EGAGVDFDGYGFCMDLSVAYVYYITDGKVRMWGNAKDAINNDFKGLATVYKNTPSFKP 74
 E G DFDG+YG+QC DL Y ++ ++ +G N+F A +Y NTP+FK
 Sbjct: 252 ENRGWDFDGSYGWQCFDLNVVYWNHLYGHGLKGYGAKDIPYANNFNSEAKIYHNTPTFKA 311

Query: 75 QLGDVAVYT---NGQYGHQCVLSGNLD---YYTCLEQNWLGSGFDGWEKATIRTHYYD 127
 + GD+ V++ G YGH VL+G+ D + L+QNW GG+ E A H Y+
 Sbjct: 312 EPGDLVVFSGRFGGGYGHTAIVLNGDYDGKLMKFQSLDQNWNNGGWRKAEVAHKVHNYE 371

Query: 128 GVTHFIRP 135
 FIRP
 Sbjct: 372 NDMIFIRP 379

>gi|2764983|emb|CAA69022.1| (Y07740) cell wall hydrolase Ply187
 [*Staphylococcus* phage 187]
 Length = 628

Score = 76.9 bits (186), Expect = 2e-13
 Identities = 50/144 (34%), Positives = 68/144 (46%), Gaps = 18/144 (12%)

Query: 5 QQAKEWIYKHEGAGVDFDGYGFCMDLSVAYVYYITDGKVRMW-----GNAKDAINNDF 59
 +Q +W G+GVD DG YG QC DL Y++ R W GNA+D +
 Sbjct: 12 KQVVDWAINLIGSGVDVDGYGRQCWDLP-NYIFN-----RYWNFKTPGNARDMAWYRY 64

Query: 60 KGLATVYKNTPSFKPQLGDAVYVYTNQY-----GHIQCVLS-GNLDYYTCLEQNWLGSGF 113
 V++NT F P+ GD+AV+T G Y GH V+ Y+ ++QNW
 Sbjct: 65 PEGFKVFRNTSDFVPKPGDIAVWTGGNWNNTWGHTGIVVGPSTKSYFYSVDQNWNNNSNS 124

Query: 114 DGWEKATIRTHYYDGVTHFIRPKF 137
 A H Y GVTHF+RP +
 Sbjct: 125 YVGSPAARIKHSHYFGVTHFVRPAY 148

309

>gi|3287732|sp|O05156|ALE1_STACP GLYCYL-GLYCINE ENDOPEPTIDASE ALE-1
 PRECURSOR >gi|1890068|dbj|BAA13069| (D86328) ALE-1
 [Staphylococcus capitis]
 Length = 362

Score = 73.4 bits (177), Expect = 2e-12
 Identities = 47/117 (40%), Positives = 61/117 (51%), Gaps = 10/117 (8%)

Query: 132 FIRPKFSGSNSKALETSKVNTFGKWKRNQYGYRNENGTFTCGFLPIFARVGSPKLSEP 191
 F++ GSNS TS N G +K N+YGT Y++E+ +FT I R+ P S P
 Sbjct: 252 FLKSAGYGSNS---TSSNNNG-YKTNKYGTLYKSESASFTAN-TDIITRLTGPFRRSMP 305

Query: 192 NGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNGKTGNSYSVGIPWG 247
 + Y+EV DG+VW+GYN G R YLPVR WN TG +G WG
 Sbjct: 306 QSGVLRKGLTIKYDEVKQDGHVWVGYNVNSGKRVYLPVRTWNESTG---ELGPLWG 359

>gi|79926|pir||A25881 lysostaphin precursor - Staphylococcus
 simulans >gi|153047 (M15686) lysostaphin (ttg start
 codon) [Staphylococcus simulans]
 Length = 389

Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)

Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK-----WKRNQYGYRNENGTFTCG 175
 HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
 Sbjct: 258 HFQRMVNSFSNSTAQDPMPLKSAGYKAGGTVTPTNTGWKTNKYGTLYKSESASFTPN 317

Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNG 234
 I R P S P + Y+EV DG+VW+GY G R YLPVR WN
 Sbjct: 318 -TDIITRTTGPFRRSMPQSGVLKAGQTIHYDEVKQDGHVWVGTYTNSGQRIYLPVRTWNK 376

Query: 235 KTGNSYSVGIPWG 247
 T ++G+ WG
 Sbjct: 377 STN---TLGVLWG 386

>gi|126496|sp|P10548|LSTP_STAST LYSOSTAPHIN PRECURSOR
 (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi|79927|pir||S01079
 lysostaphin precursor - Staphylococcus simulans bv.
 staphylolyticus >gi|581744|emb|CAA29494| (X06121)
 lysostaphin (AA 1-480) [Staphylococcus simulans bv.
 staphylolyticus]
 Length = 480

Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)

Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK-----WKRNQYGYRNENGTFTCG 175
 HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
 Sbjct: 349 HFQRMVNSFSNSTAQDPMPLKSAGYKAGGTVTPTNTGWKTNKYGTLYKSESASFTPN 408

Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNG 234
 I R P S P + Y+EV DG+VW+GY G R YLPVR WN
 Sbjct: 409 -TDIITRTTGPFRRSMPQSGVLKAGQTIHYDEVKQDGHVWVGTYTNSGQRIYLPVRTWNK 467

Query: 235 KTGNSYSVGIPWG 247
 T ++G+ WG
 Sbjct: 468 STN---TLGVLWG 477

>gi|3287967|sp|P10547|LSTP_STASI LYSOSTAPHIN PRECURSOR
 (GLYCYL-GLYCINE ENDOPEPTIDASE) >gi|2072411 (U66883)
 lysostaphin [Staphylococcus simulans]
 Length = 493

Score = 69.5 bits (167), Expect = 3e-11
 Identities = 48/133 (36%), Positives = 62/133 (46%), Gaps = 20/133 (15%)

Query: 131 HFIRPKFSGSNSKALETS---KVNTFGK-----WKRNQYGYRNENGTFTCG 175
 HF R S SNS A + K +GK WK N+YGT Y++E+ +FT
 Sbjct: 362 HFQRMVNSFSNSTAQDPMPLKSAGYKAGGTVTPTNTGWKTNKYGTLYKSESASFTPN 421

Query: 176 FLPIFARVGSPKLSEPNGYWFQPNGYTPYNEVCLSDGYVWIGYNW-QGTRYLPPVRQWNG 234

310

I R P S P + Y+EV DG+VW+GY G R YLPVR WN
 Sbjct: 422 -TDIITRTTGPFRSMPQSGVLKAGQTIHYDEVMKQDGHVWVGTYTGNSSGQRIYLPVRTWNK 480
 Query: 235 KTGNSYSVGIPWG 247
 T ++G+ WG
 Sbjct: 481 STN---TLGVWLG 490

>gi|3341932|dbj|BAA31898.1| (AB009866) amidase (peptidoglycan
 hydrolase) [bacteriophage phi PVL]
 Length = 484

Score = 68.3 bits (164), Expect = 6e-11
 Identities = 52/150 (34%), Positives = 71/150 (46%), Gaps = 17/150 (11%)

Query: 3 SQQQAKEWIYKHEGAGVDFDGYGFQCMDSVAIVVYITDGKVRMWGNAKDAINNDFKGL 62
 ++ QA++W G + D YGFQC D + + + I G+ R+ G I D K
 Sbjct: 4 TKNQAQKWFDSLGLKQFNPDLFYGFQCYDASMF-FMIATGE-RLQGLYAYNIPFDNKAR 61
 Query: 63 ATVY----KNTPSFKPQLGDVAVYTN---GQYGHICVLSGNLDYITCLEQNWLGGGF-- 113
 Y KN SF PQ D+ V+ + G GH++ V S NL+ +T QNW G G+
 Sbjct: 62 IEKYGQIIKNYDSFLPQKLDIVVFPSPKYGGAGHVEIVESANLATFTSFGQNWNGKGTW 121
 Query: 114 ----DGW--EKATIRTHYYDGVTHFIRPKF 137
 GW E T HYYD +FIR F
 Sbjct: 122 GVAQPGWGPETVTRHVHYDDPMYFIRLNF 151

Query= pt|110882 44AHJDORF012 Phage 44AHJD ORF |8391-8813|3 1
 (140 letters)

>gi|140528|sp|P24811|YQXH_BACSU HYPOTHETICAL 15.7 KD PROTEIN IN
 SPOIIIC-CWLA INTERGENIC REGION (ORF2)
 >gi|322189|pir||B44816 orf2 5'of autolytic amidase -
 Bacillus subtilis >gi|142801 (M59232) open reading frame
 2 [Bacillus subtilis] >gi|1217874|dbj|BAA06959| (D32216)
 ORF121 [Bacillus subtilis] >gi|1303767|dbj|BAA12423|
 (D84432) Yqdd [Bacillus subtilis]
 >gi|2635036|emb|CAB14532| (Z99117) alternate gene name:
 yqdd; similar to holin [Bacillus subtilis]
 Length = 140

Score = 80.4 bits (195), Expect = 6e-15
 Identities = 45/130 (34%), Positives = 67/130 (50%), Gaps = 3/130 (2%)

Query: 4 VKFRFTDSEAFHMFYAGDLKLLYFLFVLMFVDIITGISKAIKNNLWSKKSMRGFSKXX 63
 + F D ++P G +K L L VL +D++TG+ KA K L S+ + G+ +K
 Sbjct: 8 INFETLDLARVYLF---GGVKYLDLLVLVLSIIDVLTGVKAWKFKLRSRSWFGYVRKL 64
 Query: 64 XXXXXXXXXXXXXXXXXXXXKGGLLMITIFYIANEGLSIVENCAEMDVLVPEQIKDKLRVI 123
 G L T+ +YIANEGLSI EN A++ V +P I D+L+ I
 Sbjct: 65 LNFFAVILANVIDTVLNLNGVLTFGTVLFYIANEGLSITENLAQIGVKIPSSITDRLQTI 124
 Query: 124 KNDTEKSDNN 133
 +N+ E+S NN
 Sbjct: 125 ENEKEQSKNN 134

>gi|4126631|dbj|BAA36651.1| (AB016282) ORF45 [bacteriophage phi-105]
 Length = 135

Score = 76.1 bits (184), Expect = 1e-13
 Identities = 44/115 (38%), Positives = 61/115 (52%), Gaps = 4/115 (3%)

Query: 21 GDLKLLYFLFVLMFVDIITGISKAIKNNLWSKKSMRGFSKXXXXXXXXXXXXXXXXXXXX 80
 G++K L + VL +DIITG+ KA K L S+ + G+ +K
 Sbjct: 17 GEVKYLDLMLVLNIIDIITGVKAWKFKELRSRSWFGYVRKNLSFLVVIVANAIDTIMD 76
 Query: 81 XKGGLLMITIFYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKND----TEKSD 131
 G L T+ +YIANEGLSI EN A++ V +P I D+L VI++D TEK D
 Sbjct: 77 LNGVLTFTVLFYIANEGLSITENLAQIGVKIPAVITDRLHVIESDNDQKTEKDD 131

>gi|141088|sp|P26835|YNGD_CLOPE HYPOTHETICAL 14.9 KD PROTEIN IN NAGH
 3'REGION (ORFD) >gi|1075967|pir||S43905 hypothetical
 protein D - Clostridium perfringens >gi|455154 (M81878)

311

ORF D [Clostridium perfringens]
Length = 132

Score = 60.9 bits (145), Expect = 4e-09
Identities = 38/127 (29%), Positives = 63/127 (48%), Gaps = 3/127 (2%)

Query: 1 MNEVKFRFTDSEAFHMFYI-AGDLKLLYFLFVLMFVDIITGSKAIKNNLWSKKS MRGF 59
+N +K+ +I+ A D+ L+ L V +F+D +TG+ K K+ L S +RG
Sbjct: 5 INYIKWGIIVSLGTLFTWIFGAWDIPLITLL-VFIFLDYLTGVIKGCKSKELCSNIGLRGI 63

Query: 60 SKKXXXXXXXXXXXXXXXXXXXXXKGGLLMITI-FYYIANEGLSIVENCAEMDVLVPEQIKD 118
+KK + I ++YI NEG+SI+ENCA + V +PE++K
Sbjct: 64 TKKGLILVVLLVAVMLDRLLDNGTMMFRTLIAFYIMNEGISENCAALGVPIPEKLKQ 123

Query: 119 KLRVIKN 125
L+ + N
Sbjct: 124 ALKQLNN 130

>gi|2293160 (AF008220) YtkC [Bacillus subtilis]
>gi|2635548|emb|CAB15042| (Z99119) similar to autolytic
amidase [Bacillus subtilis]
Length = 134

Score = 36.4 bits (82), Expect = 0.099
Identities = 25/109 (22%), Positives = 41/109 (36%)

Query: 17 FIYAGDLKLLYFLFVLMFVDIITGSKAIKNNLWSKKS MRGFSKXXXXXXXXXXXXXXXXX 76
F + G L LM ++ I+ K + L KK KK
Sbjct: 20 PFFGGFQYSFLILLSLMAIEFISTTLKETIIHKLSFKKV FARLVKKLVTLALISVCHFFD 79

Query: 77 XXXXXKGGLLMITIFYYIANEGLSIVENCAEMDVLVPEQIKDKLRVIKN 125
+G + + I +YI E + IV + + + VP+ + D L +KN
Sbjct: 80 QLLNTQGSIRDLAIMFYILYESVQIVVTASSLGIFVPQMLVDLLET LKN 128

>gi|1181973|emb|CAA87743.1| (Z47794) holin protein [Bacteriophage
CP-1]
Length = 134

Score = 31.3 bits (69), Expect = 3.3
Identities = 27/88 (30%), Positives = 36/88 (40%), Gaps = 5/88 (5%)

Query: 29 LFLVMFVDIITGSKAIKNNLWSKKS MRGFSKXXXXXXXXXXXXXXXXXXXXXK--GGLL 86
LF L+ D ITG KA K S ++G K G +L
Sbjct: 18 LFAILLPDFITGFLKAWKNKVTDSWTGLKGVIKHTLTFIFYFVAVFLTYIHAMAVGQIL 77

Query: 87 MITIFYYIANEGLSIVENCAEMDVLVPE 114
++ I Y A LSI+EN A M V +P+
Sbjct: 78 LVIINLYYA---LSIMENLAVMGVFIPK 102

Table 21

Phage 182 complete genome sequence. 17833 nucleotides.

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1      tagaatattg tcataaaaca caaacataat aatgcatatt attgtttaca aatatgtaat ttcgtgatat
71     aatatatttg taagttaaag gaggtgacaa aagaacaaat cataaatgct ttagaaattg caaaaactat
141    tggaggaaaaa ataatagaat attcactaca acaaatagat gaaattaaat caacaatttt cagaattaga
211    ttaaaaaggc atgaactaga ggaattggcg gacgaagtaa acgatattgc taaagatccg gaggaaagat
281    atctttttatc gttttattac acagaagaag aacggttggc tgaaattccc tctgcaagat taatagatta
351    ttacaacgaa aagatcacaa atctgaaatc ggaaatcata tcaactcgaa aaagattaca aaaactagta
421    aaataaattac acaaaaagct ttacaaatat aacacatcat gttatactaa aagagtagta agggaaacgga
491    aaatacctta cttcacacct caatcattct tatcaaaata caaaaggagg gaaaataatg ggtcgaaaac
561    taatgcaacg aaacgtaaca tcaactaaag tagaattctc agaagttatc gtacaagatg gagcgccaac
631    aattgtacca tgcgaaccag ttgtcttaac aggaaaaactt tcagaagaaa aagctttatc agcgatcaaa
701    cgtaaaaaac ctgataaaaa cgtagtgtga acaaatgttt cacatgaaac agcgctttac acaatgccag
771    tcgataaatt tatcgagtta gcagacaaat caacacaaag ctaataaaaa caaaactaaa acaaaaacaga
841    ggagattata atcatggaaa tcgtaaaaaa cacatttgac acacaaacac cagaaggaaat gttacaagta
911    ttcaatgcca caaacggggc ttcaattccg ttacgtaacg caattggcga agtactagaa ttgaaagata
981    ttctagttta ctcagacgaa gtttctgggt ttggtggagc cgaaccatca caagcagaac tagtctgctt
1051   cttcacagaa gatggtaaaa cttatgcccc tgatcatcaa ccaaaaattt ctttctgcta aggaaaaatc aacggtggac
1121   gatattgatga ctgctaaccg tgacatcaaa ccaaaaattt ctttctgcta aggaaaaatc aacggtggac
1191   aaaaaattgt aaatctacaa gtggtttcac tgtagcataa aaatacagga atctagtaag ccacttagcg
1261   aatctcgcta ggtggttttt attatgtttc tacattgagg tgtgtagaat tgaccgtaag aatatcaaa
1331   aatgatagag ccaagttaga gaaaatctac ggtaaatcta acaaaagctc taaaaaatc aatcgtttaa
1401   gacaaaaagg agttgaggaa aggcaacttc caactgttcc aacatcaaa aaaagactta ttgactacgt
1471   aaaaatcaaca aatatgagtc gtatgtatgt taacaagatg ttagacgagt tggtagattt tgcacaacct
1541   taacacgaga attacatttt tgagatcaac aagcgaaatg ttgcaatctc aagagcgcaa atcaaaagaag
1611   cgcaaatata aacagagcaa gctcaaaaag cgaaagaaga acactacaaa gagcttaaca aagtgaagt
1681   taagaagccc acagaaaaca caattgtcac accaactatt ttaacagagt taggtgctga cttacctttt
1751   caagcaatac cagattttta tattgacgct ttcacttctc cagaaggagt tcagtcttat ttgaaaaata
1821   taggaaaaaca agacgaacaa tattttgacg aaagagacca actttattac gacaatttca gacaagcgat
1891   gtttactatt ttcaattcag acgctgacga tattgttcgt ttacttgact caatggggct tgatctattt
1961   atgaaaacat atggttagta cttcttagac atgaaccttg actacattta tgacgaagca gaagtacaac
2031   agaaaaaaga acaagtttac agtaagattg caaaagatg cgagtctgaa acaggtggag aagtcacctc
2101   atataacccc acgaagaaca tcacaattaa ttcagaaaca ggagaagaat tatgattaag aaatatactg
2171   gcgactttga aacaacaact gatctcaacg attgtcgtgt atggctcgtg ggcgtatgag atatagacaa
2241   cgttgacaa atgacggtcg gtttagaaat cgattctttt tttgagtggg gtataatgca aggcagcaca
2311   gacattttat tccacaacga aaaatttgac ggagagttta tgccttcatg gttattcaaa aatgggttca
2381   aatgggtgtaa agaagcaaaa gaagatcgaa cattctccac actcatatca aatatgggtc aatgggtatg
2451   tttggaaaatt tgttgggaag ttaattcac aacaacaaaa tcaggttaaaa cgaaaaaaga gaaatctcga
2521   acaataattt atgatagcct taaaaaatat ccttttcag tgaacaaaat tgcagaagct ttaattttc
2591   ctataaaaaa aggcgaataa gattatacaa aagaagacc tattgggtac aaaccaacaa aagatgaatg
2661   ggagtattta aagaacgaca ttcagattat ggcgatggca ttaaaaattc aattcgatca aggactaact
2731   cgaatgacta gaggaagcga cgcttttagc gattacaaag attggctaaa agctacacat ggaaaaatcaa
2801   ctttcaaaaca atggtttcct attttgtctt tagggtttga taaagactta cgtaagcat acaaaaggcg
2871   cttcacttgg gtaaacaaag tttttcaagg gaaagaaata ggtgacggca ttgtcttga tgtcaactct
2941   ttgtatccct ctcaaatgta cgtaagacct ttaccatag taacacctct attctacgaa ggagaataca
3011   aaccgaacaa cgactatccg ctgtacattc aaaatatcaa agtaagattc cgtttaaaag aggggtatat
3081   tccaaccatt caagttaagc aaagttcatt attcattcaa aacgaatata ttgaatcaag tgaatacaag
3151   ttaggagttg acgaattaat cgactcttact cttacaaatg ttgacctaga attatttttt gaacactacg
3221   atattttaga gatacattac acttacggat atagtgtcaa agcttcttgt gatattgtca aaggctggat
3291   cgataaatgg atcgaaagta agaaccaccac cgaaggggct agaaaagcta acgccaaggg tatgttaaat
3361   agcttgatg gaaaagttcg aacaaaccct gacattacag gaaaagtgcc ttacatgggc gaggaacgca
3431   ttgttcgatt gacactagga gaagaagaat taagagatcc tgtttatgtt cggcttgcta gttttgtgac
3501   ggcttggggg agatatacta ccatatacaac cgctcaaaaa tgttttgatc gcattattta ttgtgatata
3571   gtagcattc atctagtagg aacagaagtc ccagaagcaa tgcgactctt ggttgatctc aaaaaacttg
3641   gttattgggg gcatgaaagc acattttcaac gagcaaaatt cattcggcag aaaacatacg tagaagaat
3711   tgatggcgaa ttaaatgtaa agtgtgctgg tatgccagat cgaataaaaag agattgtaac ttttgacaat
3781   tttgaagttg gtttttcaag ctatggaaag ttgtactcta aaagaacaca aggtggcggt gtattagtag
3851   acacaactgt tacaatcaaa taaggaggac taataatgga actatataaa gcaatgttta tctgactgga
3921   tgaaggtact attgacggtt acgatactga acactatgta gatatttctt tacatgactt tgaagaaata
3991   tatggaaaaa aaacacgtga aattgaagca gtaacattag taaaaacagg aaatttaaaa aataaatta
4061   tttacatcct ttgcaagata ttgttaaaata tcttctgtat agttgacaag agtcaaaattt ggcagagattg
4131   ggcgaatgta cagtgaaat atcgtgcgct cccgttaagt tatggacaca taaacgtttt gaccgtcaac
4201   caatcgcaaa aaccttttag gtagtgcctt taaatgtggc tactcttttt tgtgtttcac agaattatgt
4271   ttcaactgaa acagttttta tggtaataa gaatcaaaa gaggtggaga ttatggaaat taaagaacct
4341   gaatacaatt taaatggtat tcttgaaagt gtcacagacg gtgaagcaag atcaaaagatt gtagaacatc
4411   ttgaagcatt gcgagaagac tacggagcaa caactgaagc tttgacatca gcaaatagca cacttgaaaa
4481   gttaaagaaa gataacgaag cgttggttat ttcaaaactc aaattgttcc gagaacgagc gatcgtagaa
4551   ccagcagaaa ataacgaacc agaaacgagc cagaatatta cactagacga tttaggaatt taaggaggaa
4621   aaaacatggc tgacaaaatc acagaacaag atgttcttcg tgccacaaat gtagaacacac cagtacaatt
4691   aatgactgct atttataata gttcatcatc tctttttcag gcgaacgtac ctatgccaaa tgcagataac

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4761 atcgaagcgg ttggtgcagg gatcacacgt ttagacgtag taaaaaacga atttatttca actttagttg
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 7141 caagatgtta atcgataatg atcctaacga ttagggaggt aaatctgact atgcttctgt attcatgcaa
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 7351 catcatggga gcaggaacac aagtaaacaa ctatgtttct gaaaaagaaa acgggttgaa cctcttggca
 7421 ggtaaatggt cagatatcga aaatattcca gataatgtaa cacagcttgg atcaaaacta cttttcaca
 7491 caggaaactt tcaaaactat tatcaattgc gcttcaaa aattaaatat gagtatgcaa caagacttga
 7561 tcgttacttc tcaatgtatg gcacaaagag caatcgagta gctacaccaa acttacaac aagaaagca
 7631 tggaaatttc ttaaattaaa agaaccaaat attgtaggca caatgagtaa cgtatgatta acacgtgtga
 7701 aacaaatttt tagtgcaggc gttacgcttt ggcatacgaa tgatgttttg aattataacc aagacaacgg
 7771 agatgtatag gaaggaggaa taagatgagt agacgaaaag gtgcaggact tgctagaat aaccgttata
 7841 cagcaaaaag cagaccttat ccaaatgaac cctattcaag tgatgtagaa gaaatcagct actatgaaca
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10081	gcaatcaata	aagttagtgg	ctggaatacc	gctacaggag	atattttatct	taacattaaa	ggaacggagg
10151	gtgtataatg	gcagacatta	gaacacaact	aacaagtga	gatggatcag	acaatttatt	tccaatttca
10221	aaagccggtt	atattatgac	taatagcggt	acgaatgtag	aaggagaatt	gggtacactc	aaacaaaatg
10291	acgaaacaat	gaatacctca	gttcaaaatg	ctgtagttag	tgccaatcaa	gcaaaagatt	ctgtagctga
10361	attaaatgta	aatgttggta	aactaaccaa	tcgaataaca	acattagaga	gtacagtggc	taatcttgat
10431	gggtattcgtt	atgtagaggt	gtaatatggc	agataaaaa	attcaaatgc	aggataaaga	tcataatcgt
10501	ttaatgctcg	ttacaattgc	taaaaatggt	ctaacaggcg	actctaactc	tgaattagtt	aatgctgaaa
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10641	agaaattgac	acagtacat	caaccgcaa	tcaagcgttg	acgaaggctg	gtacagcaca	acaaaccgca
10711	gaacaagcga	aaacaacagc	aaacagtatc	agcgcagttg	caacggcagc	taaaaacaca	gctgattcag
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10851	attaccatag	gaggaaaaat	aatggcaaat	aaaaatattc	aaatgaagga	tagcaatgac	aataatttat
10921	atccaaagtg	tcgagcagaa	aacttggttag	atttgaccag	tcgtgctgaa	ttaacaatga	caaatgttca
10991	attatctgca	gctggtgata	aaacaaatgc	aactctttat	ctcgggtgca	taggtatgct	cgaaggtagt
11061	ataaagttta	ctgaaagttt	gacaaacctt	gtgatcaca	cgctaccaga	agggtttaga	ccaataagaa
11131	caaaacgtat	tggttggttc	gcaaaatatt	acacaccaa	tccaacagat	acaaaagaaa	tggtttatgt
11201	atcaatcaca	cctgatggca	aagtaactgt	aatgacaat	gtaggtaaaa	tcgaatatct	atccctagat
11271	aattgcggtt	tccctctaaa	ataaggaggt	tcataatgaa	gaacgaattg	atattcaaat	gaacaagatg
11341	aaagaagaaa	atcaaaagaa	ttacattatg	cacctgaaa	cgaacccgaa	acaagttggt	tttgatgaaa
11411	cattgcatgg	aaatgaaaat	caggagaggt	tcaacaattt	tggtgacaca	agaaaaatga	caactacaat
11481	tgatgtaagt	gcttatgggg	gtatcgctga	cggtgtaaca	gattgtacac	caatatataa	taactactct
11551	gaagaaaaaa	gcgaatggg	tatcactttt	tatttctctc	cttgtagaacg	tgattcatat	tatcgctttg
11621	ctaaccacct	tgaattgaaa	cgtagatgac	ctgtagttag	tttcttagga	tcgggagaaa	cgacattaaa
11691	gtttgaaaca	atgacggcat	ttaatgtaaa	catcgaaagt	ttcaaatattg	atgggtttgc	attatgggtg
11761	ccacaaggcg	ctcaaagtgg	taaagggaat	ttctttaatg	atactcgcaa	ttacaatcgt	tttgactttg
11831	atttggttgt	tcgttaactgt	actttaaatg	aaggaaacgt	tggtgtgtgt	gctagaggta	gagggtttac
11901	atttgaataa	tgctattctt	ctaatacttc	tcaagcaatt	atcaaaacag	cttttcccca	gttaaatggt
11971	atgtggcaag	ggaacgatat	caataactagg	ggtacagggt	ttagagggtt	ctttgtgaaa	aacaaccgta
12041	ttcatttttg	tacagcgatc	attatcgaca	atgacgatga	ttatcagaat	gtaattaatt	ttctgtgaaat
12111	ttctggtaac	acaatcgaag	gtggcgtaag	ttattatcga	ggatatgcgc	ataacttgca	tgccaaaac
12181	aacaaccatt	ttctagcata	cggaaataga	aacgctttgt	ttgagtttca	agatgtggat	caagcttata
12251	ttgatgtaga	tggtttattgt	cgtaactcac	aagtcgaggg	aatgaatagt	acagctatct	caagtttaat
12321	tggtgtttac	ggacattacc	gaaacttaaa	gattacaggt	aaattatctc	gttgctcaag	acatgtttac
12391	acgttgtagc	gcggtggcgt	taatttctat	tgtagcttga	tggcacaaga	agcacctttg	acggacgggt
12461	accggtttat	tcaaacggct	gacaatcgag	ttaactatga	tggtgtttgt	gttcgtggtt	tgcttaattc
12531	aacaaaagta	aatacaccaa	tgatctataa	agcacctcag	actgttttct	ataatcgtag	aatcgatcat
12601	gtgctaaacg	gtccaaatgc	aagtaaatga	tataactagg	aggatatgag	atggcaactc	ttacaatatga
12671	acaaatagct	agaggacaaa	caatcgctaa	aatactttca	aaatatggct	ataataaaaa	ttcacaaagta
12741	ggagttgtcg	ccaatctcca	ttgggaatcg	gctggtttga	acccgaacag	caatgaatat	ggtggaggcg
12811	gatatgggtt	aggtcaatgg	acgcctaaaa	gcaatcttta	tcgccaagca	caaatttggt	ggttgctctaa
12881	tgctaaagct	gaaacggttg	aaggtcaagc	agagatcatc	gctcaagggg	ataaaacagg	tcaatggatg
12951	gataatacac	ctgtttcttc	tgtaggttat	actaaccttc	agacctttc	agcatttaaa	caatctgcaa
13021	atattgtagt	tgctacaatt	aattttatgt	gtcactggga	acgacctggt	aaactcata	tcgaagaag
13091	acttgatctt	gcacaagctt	atagtaagca	tattgacggt	agcgttggtg	gtggcgtaaa	acgttgctat
13161	ggaaccccaa	tcaagaatac	aaatcttgat	cctaaaagtt	tcatgagtgg	acaacttttt	ggcacgcatg
13231	caggaaacgg	cagaccaaat	aatttccatg	atgggtttga	ctttgggttca	attgatcacc	ctggcaatga
13301	aatgtgggca	tggtgcgatg	gaacagtaac	acatgtttga	acaatgggag	cattaagagc	gtattttgtg
13371	ataaatgatg	gtacttacaa	tatcgtttat	caagaattta	gttataacca	gtcaaatata	aaggtaaaag
13441	ttggcgacaa	agtttaagaac	ggacaagttt	gcgcaatacg	tgacgcggat	cattttacat	taggttttac
13511	taaaaaagat	tttatgactg	cggttaggac	ttcttctata	gatgatggaa	catgggaaga	cccttggaag
13581	tttttagggc	aatgttttgg	agatggagat	actggcggag	ataatgacga	taacaataag	gataaaaatg
13651	atcttatatta	tctattgcta	tccgatgcct	tgaatgggtg	gaaattttta	taaggagaaa	aaggtagtat
13721	agaatatcta	acacaatggg	tgccagatga	tatcatcttt	gtttatgggt	tgatttatgt	tgatttatgt
13791	gcaatgatta	tcgattttgt	gttaggtttt	acaattgcca	aatttaacaa	ggaaatcgac	tttagtagtt
13861	ttaaagctaa	agcagggtatc	attgtttaagg	tgccagaaat	gggttttagtg	gtttacttta	ttctgtagc
13931	agtaaaattc	ggtgcagtag	gtattacaat	gtatataaca	atggtgggtg	gtttgatttt	atcagaatt
14001	tatagtatac	taggacatat	ttcagatata	gatgatgata	ataattggac	tgatttatgt	aagaagtttt
14071	tagacggaac	actcaacaga	aaggacgata	ttaaatgatg	aatggtattg	atatctctag	ttatcaaaaa
14141	ggaattgatc	tttcaaaagt	tccatgcgat	tttgtaaaata	ttaaagcaac	aggcgggaaca	ggttatgtaa
14211	accctgatgt	tgaccgagca	tttcaacaag	tttgcctttt	aggtaaaaaag	attggttggtg	atcattttgc
14281	gcattgagagg	gggtttagaag	gtacacctca	acaagaagcg	caattctttt	tagataatat	taagggttac
14351	attggtaagg	ctgttcttat	tcttgacttt	gaagggtcaa	atcagaaaaga	tgtaaatgtg	gcgaaagcat
14421	ttcttgattta	tggtttataat	aaaaacaggc	tttaaacgatg	gttttatacag	tatacagcaa	acctcaatat
14491	aactgatttt	tctagtattg	caaaaggcga	ttatgggtta	tggtgtgctg	aatatggatc	aatcaacca
14561	caaggctact	ctcaaccagc	gccacctaaa	acaaataatt	ttccaattgt	tgccgttttt	cagtttacaa
14631	gtaaaagacg	tttaccagga	tacaacggca	actctgattt	gaattgtttt	tatggcgatg	gttaaatcatg
14701	ggatctgtat	gtaggtaaaa	aacaggatca	aattgttcct	cctgaaaata	aaatatttga	cgccacaagt
14771	gatgagttta	ttttcactct	tacaacaggt	agcacaagcg	tggtttattt	tgacggagaa	acgatctttg
14841	aattgtctga	tccaacacaa	ctcgatcata	ttagaggaaac	atacaatcat	gttcatggaa	aagaaatccc
14911	atcaatgggtg	tggaacacctg	aacaatttga	tatttactta	aaaatgtatg	aaaagaaacc	agtatataag
14981	taggagtgta	tagtatgaca	aatagcttag	gcgttaaaact	tgaagagaaa	aacttatact	ataacacctaa
15051	caatgcttta	ggttttaatt	gcctaatggt	gtttgtaata	ggcgacgctg	gtataggtaa	aaacttatggt
15121	tataaaaaat	ttgttggtta	tcgcttttat	aaacacggcg	aaacaatttat	ttatttaaga	agattcaaaa
15191	cagaacttaa	aaagattcct	caatttttca	aaacaatggc	gaaagaatttc	cctgatcata	aacttgaagt
15261	aaaaggaaaa	gaattctatt	gtgatgataa	attaatgggt	tggtgctgtt	cacttagtac	gtggggaatt
15331	gaaaaatcta	atgaatatcc	cgaagttcgt	caaattttgt	ttgatgagtt	tttaattgag	aaatcaaaaa

15401	tcacttattt	accaaacgaa	gctgaagcct	tattgaacat	gatggaaacg	gttttccgaa	gacgtacaaa
15471	tacaagatgt	gttatgttga	gtaatgcaac	tagtgtagt	aacccttatt	tcttgtat	caatctgcag
15541	ccagatttga	ataagcggtt	taatctatat	caagatcgag	gtatattgat	tgaattgtgt	gattcaaaag
15611	accttgacga	agtgaagaga	gaaacacctt	ttggtagatt	gattcgtgga	acagaatacg	aagattttag
15681	tatcaacaat	gagtttgtca	atgatagtga	tacgtttatt	gaaaagagaa	gtaaaaatag	tagtttctta
15751	tcgcccattg	cttttgaagg	gaaaatcttt	gggtattgga	tagacgctga	aacaggttgt	gtctatgtga
15821	gttatgatta	tcaaccaaat	acaaatcatt	tttatgcaat	gactacgaaa	gaccatgaag	aaaatagatt
15891	gctgatgaaa	aattggcgaa	ataattatta	tctttcaaca	gtggcgaaa	cattcaagaa	tagttatctg
15961	cgggttgata	acattgttat	taagaattta	cattatgatt	tgtttaataa	gatgaaatc	tggttaacct
16031	atttttagtag	agctaccacg	attagttcta	ttacaatgat	gaatagtaga	taacatagta	attgtagtct
16101	gcgatagttt	tgttttgggt	ctttggcggt	agtgattttt	gctaacgcct	ttttgtttgc	ttttggatcg
16171	gggtgtgtaa	tgtagacgaa	atctttcttc	atagtctctt	ctccttatac	agttttaata	attccctgta
16241	aaatgtagct	ataggacgct	catttctttc	tattctaacy	caattcacta	tatccatttc	taggtatata
16311	cggctatatt	ttaatgcttt	tgtaaagggt	agaggttcgg	ttttgtgtat	caaaacctcc	caaccatcta
16381	tataaaatc	tgtagatcgc	tatatgggt	ccttgttagaa	tgtagccatt	attccacctc	ctttaaatag
16451	ccttttggtta	tttgtaacgc	taactgatag	cgagaaccaa	cttttacgta	tgaagtact	aatttcattg
16521	cctgacaata	cttttcaaga	atgttaaatt	gactcgattc	gggtaatagc	gttgaatgag	ttaacaaaag
16591	ttcggtgata	tttatttccg	gaacgtcgaa	atcttgtaaa	gtccctcta	tgatctctat	tttttcattg
16661	tctgaaaggt	tacgtttaca	gtagaacgt	aaccattcaa	ttagtccgcg	gtgttctttg	aatgttcgtg
16731	caatcatttt	aattcctcct	atttgcctgt	aatttggtta	tatccgtcat	gtttcaattg	ttccgcatag
16801	tggtcaacgc	ttttcattga	tttcgttatt	gcgatattaa	tgcaatggct	atcaagataa	acatagtatt
16871	atttatcatg	tgtaaacacg	aactcttttg	taacgtaatc	aatgtataaa	attaattggt	ttcctccttg
16941	tgttatttct	gacttgatag	acgctaaact	atcgttgctc	tctttagtta	gttgatttaa	accctctaaa
17011	attaatgata	aattgttaat	catgtaaaac	actcctttta	tattaatttg	atattgatac	caccaatcga
17081	ataagattgg	tagcattgta	tcgaattaat	atgttatttc	tgtagttttc	catgaatact	cggaataaag
17151	atccatattc	aattccttta	gttcttcaaa	agataacaaa	caatattcct	catcgcttac	ctcatcaata
17221	tcaataagat	aatgtttatt	gttttcggta	tctatgatat	gataattcat	atccactca	ttaaaggggt
17291	gaagtagaga	tacctctcct	ttttcagcta	ttaatgattt	attgttcata	tgaaacactc	cttttatatt
17361	aatttgatat	tgataccacc	aatcaaatgt	gattggtagc	attgtattaa	attaatattc	tggaataatt
17431	attgagaaa	tccagttatc	atcaaatgaa	attgttttat	tttcaagtaa	ctttttagcc	tcatccacct
17501	caaattctaa	atagaggaat	ttactaagtt	tatcctcatc	tctaaaaatt	ttcatacata	ccacgttatt
17571	tgaataaatt	tctgtgtata	cgatcgggtc	attcatgttt	atcatccttt	ctttattaca	tatatagtat
17641	atcatgtatt	tacatatatg	tcaatcattt	aattcattta	ttttaatgat	ttatttgatt	gtttttttat
17711	gatcctttct	ttattacatc	tattattatg	catgtatgat	tgattttgtc	aacaattaaa	ttcatataaa
17781	tgtagtttgg	ggtcagttac	atttgtgtta	tcaaaaaaag	ataatattct	att	

Table 22

Phage 182 ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	182ORF001	2	5966..7780	604	Tail protein;
2	182ORF002	1	2152..3873	573	DNA polymerase;
3	182ORF003	1	11305..12639	444	
4	182ORF004	3	4626..5954	442	Major head protein;
5	182ORF005	3	12651..13700	349	Glycyl-Glycine endopeptidase; Lysostaphin precursor;
6	182ORF006	1	14995..16026	343	Encapsidation protein; ATG/GTP-binding site motif A;
7	182ORF007	1	7795..8775	326	Upper collar protein;
8	182ORF008	2	14105..14983	292	Lysozyme; Muramidase;
9	182ORF010	2	1310..2155	281	Terminal protein;
10	182ORF009	2	8765..9601	278	Lower collar protein;
11	182ORF011	1	9607..10158	183	Pre-neck appendage protein;
12	182ORF012	3	10872..11294	140	
13	182ORF013	1	10456..10860	134	
14	182ORF014	3	13716..14108	130	Lysis protein;
15	182ORF015	2	854..1225	123	Early protein;
16	182ORF018	-2	16429..16737	102	
17	182ORF020	3	10158..10454	98	Leucine-zipper motif;
18	182ORF019	3	4323..4613	96	Head protein;
19	182ORF016	-3	16749..17033	94	
20	182ORF022	1	12868..13149	93	
21	182ORF023	-2	11914..12189	91	
22	182ORF017	1	154..426	90	
23	182ORF024	3	6174..6446	90	
24	182ORF025	2	548..814	88	Early protein;
25	182ORF026	-3	12999..13259	86	
26	182ORF027	-1	14642..14896	84	
27	182ORF028	3	14430..14672	80	
28	182ORF021	-3	17106..17339	77	
29	182ORF030	-1	16199..16429	76	
30	182ORF031	-3	8379..8603	74	
31	182ORF032	-1	11195..11413	72	
32	182ORF033	-1	4727..4942	71	
33	182ORF034	-1	5951..6160	69	
34	182ORF029	-3	17412..17606	64	
35	182ORF035	-3	15570..15758	62	
36	182ORF036	-3	2127..2315	62	
37	182ORF037	-1	12095..12280	61	
38	182ORF038	3	14769..14951	60	
39	182ORF039	2	9992..10171	59	
40	182ORF040	-3	16029..16202	57	
41	182ORF041	1	3886..4056	56	Early protein;
42	182ORF042	-3	10671..10832	53	
43	182ORF043	-3	10491..10652	53	
44	182ORF044	-1	6299..6457	52	
45	182ORF045	-2	6571..6729	52	
46	182ORF046	2	2372..2527	51	
47	182ORF047	-2	13201..13353	50	
48	182ORF048	-3	3243..3395	50	
49	182ORF049	3	1578..1724	48	
50	182ORF050	2	8012..8155	47	
51	182ORF051	3	9390..9530	46	
52	182ORF052	1	4096..4233	45	
53	182ORF053	2	15656..15793	45	
54	182ORF054	-2	8002..8136	44	
55	182ORF055	2	8324..8455	43	
56	182ORF056	3	6549..6680	43	
57	182ORF057	-3	8133..8264	43	
58	182ORF058	-1	5048..5176	42	
59	182ORF059	-2	15748..15876	42	
60	182ORF060	-3	15276..15404	42	
61	182ORF061	-3	1974..2102	42	
62	182ORF062	-2	1867..1992	41	
63	182ORF063	-3	14181..14306	41	
64	182ORF064	-2	7234..7356	40	

317

65	182ORF065	-2	3460..3582	40	
66	182ORF066	1	4234..4353	39	
67	182ORF067	-1	13763..13882	39	
68	182ORF068	-1	7148..7267	39	
69	182ORF069	-3	4908..5027	39	
70	182ORF070	-3	912..1031	39	
71	182ORF071	2	11741..11857	38	
72	182ORF072	-3	11610..11723	37	
73	182ORF073	-3	2763..2876	37	
74	182ORF074	-1	8813..8923	36	
75	182ORF075	-3	7353..7463	36	
76	182ORF076	-3	2316..2426	36	
77	182ORF077	2	11858..11965	35	
78	182ORF078	-2	7564..7671	35	
79	182ORF079	-2	7381..7488	35	
80	182ORF080	-2	4372..4473	33	

Table 23

Predicted amino acid sequences of ORFs from phage 182

182ORF001

5966 atggcagaaggtatacaaatgtaaaattgttggttaacgtgccttttgataaacacctatacacacacaagatgggtttaaaact
 1 M A R R Y T N V K L L A N V P F D N T Y T H T R W F K T
 6050 caacaggaacaggaatcgactttaattcgcttctgttcttaacgagaatagagattgttcttatcaaagggatacacaaactc
 29 Q Q E Q E S Y F N S F P V L N E N R D C S Y Q R D T Q L
 6134 gggggagtttttagagtagataaaacacaaagacgcttatatgcttgaactatctcatctttaaaccgaagaaacttatcct
 57 G G V F R V D K H K D A L Y A C N Y L I F K N E E T Y P
 6218 agtaaatggcagtatgcctttgttactgatattgaatataagaatgacaacacagtttgccttacctttgaaattgatgtttta
 85 S K W Q Y A F V T D I E Y K N D N T S F V T F E I D V L
 6302 caaacttatcgcttgcgatattgggtatacagagaagtttcattgcaaaagaacaccctcaactttattattcgaatggaatacct
 113 Q T Y R F D I G I R E S F I A K E H P Q L Y Y S N G I P
 6386 ttcttaatacaattgaagatcgcttgattacggtagagaatcacacaacaacaatgtaacaacttttcatcctaagcatgga
 141 F I N T I E E S L D Y G R E Y T T T N V T T F H P N D G
 6470 gtcaattttctgttatttcaacaagtgaagcaatgccagttggagataaggaagataaatcaggaggatcaatagtaggtggc
 169 V N F L V I L T S E A M P V G D K E D K S G S I V G G
 6554 ccatctccttttcttatttttcttcttcaattcaagtggggagggtatacaaaccaaatggggagggaatgtaatttt
 197 P S P F S Y Y L L P I N S S G E V Y K P N G A G N A N F
 6638 ggagagtacatggcgtttcttacaagaaagaccttttttaataagatagtcgggatgtatgtaacgtcgtatatacagggtata
 225 G E Y M A F L T T K E P F L N K I V G M Y V T S Y T G I
 6722 ccattcattgtggatcacgcgaacaaaacggtaaggtataatgcaggaggttcttataagatcatgcttccaacctacgctagt
 253 P F I V D H A N K T V R Y N A G G S Y K I M L P T Y A S
 6806 gatccaacaggaacaatgaaaacattcgcttcttcttctgtgtaaaagaagcaagaacattcgctacctaagaattgacttctgta
 281 D P T G T M K T F A F F C V K E A R T F V P K R I D L V
 6890 gggaacgtgtataactacttttagagaagcttttccgttttaattgtaaggaatcaaaactattttatgtatcccttctgttttaata
 309 G N Y N Y P R E A F P F N V K E S K L F M Y P Y C L I
 6974 gaaattacagatacaaaaggacatgtaattgactttaagacctgaatatcttacagggtggtaaaattgagtgatatgttaaagggt
 337 E I T D T K G H V M T L R P E Y L T G G K L S V Y V K G
 7058 tcgttaggaatttctaataaagtgatgacgagcagcttatgatgtaagtaactcaaccattattaccaatttaagtgc
 365 L I S N K V M I E P I D Y D V S N S T I I T N L S D
 7142 aagatgttaatcgataatgatcctaacgatgtaggagttaaatctgactatgcttctgcattcatgcaaggaacaaaaactcc
 393 K M L I D N D P N D V G V K S D Y A S A F M Q G N K N S
 7226 ttgattgctcaagagcaaaacattcgcaatacttcagacatgggtatgggaacagtgcaatgagtaacaggagcgatcctt
 421 L I A Q E Q N I R N T F R H G M G N S A M S T G G A I F
 7310 tcagccttagcaagtaacaaccttttgggttgacttaacatcatgggagcaggacaacaagtaacaactatgtttctgaa
 449 L A S N N P F V G L T N I M G A G Q Q V N N Y V S E
 7394 aaagaaaacggtttgaacctcttggcaggtaaagtggcagatatacaaaaatttccagataatgtaacacagcttgatcaaac
 477 K E N G L N L L A G K V A D I E N I P D N V T Q L G S N
 7478 ttatcttccacaacaggaaactttcaaaactattatcaattgcttcaaaacaattaaatagagtatgcaacaagacttgat
 505 L S F T T G N F Q N Y Y Q L R F K Q I K Y E Y A T R L D
 7562 cgttacttctcaatgtatggcacaagagcaatcgatgactacacaaacttacaacaagaaagcattggaatttcattaaa
 533 R Y T S M Y G T K S N R V A T P N L Q T R K A W N Y T T
 7646 ttaaaagaaccaaatattgtaggcacaatgagtaacgatgtattaacacgtgtgaaacaaatttttagtgcaggcggttacgctt
 561 L K E P N I V G T M S N D V L T R V K Q I F S A G V T L
 7730 tggcacaagaatgatgttttgaattataaccaagacaacggagatgtatag 7780
 589 W H T N D V L N Y N Q D N G D V *

182ORF002

2152 atgattaagaaatatactggcgactttgaaacaacaactgatctcaacgattgtcgtgtatggctggtggcgatgcatata
 1 M I K K Y T G D F E T T T D L N D C R V W S W G V C D I
 2236 gacaacggttgacaatatgacgttcggttagaaatcgattcttttttgagtgggtgtaaaatgcaaggcagcacagacatttat
 29 D N V D N M T F G L E I D S F F E W C K M Q G S T D I Y
 2320 ttccacaacgaaaaatttgacggagagtttatgcttcttatggttattcaaaaatgggttcaaatggtgtaaaagaagcaaaagaa
 57 F H N E K F D G E F M L S W L F K N G F K W C K E A K E
 2404 gatcgaacattctccacactcatatcaaatatgggtcaatgggtatgctttggaaatttgggtgggaagttaattacacaacaaca
 85 D R T F S T L I S N M G Q W Y A L E I C W E V N Y T T T
 2488 aaatcaggttaaaacgaaaaagagaaatctcgaacaataatttatgatagccttaaaaaatatccttttccagtgaacaaatt
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 2572 gcagaagccttttaatttctataaaaaaggcgaaatagattatcaaaaagaagacctattgggtacaaaacaaacaaagat
 141 A E A F N F P I K K G E I D Y T K E R P I G Y K P T K D
 2656 gaatgggagttttaaagaacgacattcagattatggcgatggcattaaaaattcaattcgatcaaggactaactcgaatgact
 169 E W E Y L K N D I Q I M A M A L K I Q F D Q G L T R M T
 2740 agaggaagcgacgcttagcgattacaaagattggctaaaagctacacatggaaaatcaactttcaaaacaaatgggttcttatt
 197 R G S D A L G D Y K D W L K A T H G K S T F K Q W F P F
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 225 L G L G F D K D L R K A Y K G G F T W V N K V F Q G K E
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 253 I G D G I V F D V N S L Y P S Q M Y V R P L P Y G T P L
 2992 ttctacgaagagaatcaaaacgaaacgactacgctgtacattcaaaatatacaagtaagattccgttttaaggagggt
 281 F Y E G E Y K P N N D Y P L Y I Q N I K V R F R L K E G
 3076 tatattccaaccattcaagttaagcaaaagtctatttattcattcaaaacgaatatcttgaatcaagtgtaaacaagtttaggagtt

319

309 Y I P T I Q V K Q S S L F I Q N E Y L E S S V N K L G V

3160 gacgaattaatcgatcttactcttacaatgttgacctagaattatttttgaacactacgatatttttagagatacattacact
337 D E L I D L T L T N V D L E L F F E H Y D I L E I H Y T
3244 tacggatatatgttcaaagcttctgtgatgttcaaaggctggatcgataaatggatcgaagtaaaacaccaccgaaggg
365 Y G Y M F K A S C D M F K G W I D K W I E V K N T T E G
3328 gctagaaaagctaaccgcaaaaggtatgttaaatagcttgatggaaagttcggaacaaacctgacattacaggaaaagtgcct
393 A R K A N A K G M L N S L Y G K F G T N P D I T G K V P
3412 tacatggcgaggacggcattgttcgattgacactaggagaagaattaagagatcctgtttatgttcgcttgctagtgtt
421 Y M G E D G I V R L T L G E E E L R D P V Y V P L A S F
3496 gtgacggcttggggtagatatactaccattacaacccgctcaaaaatgttttgatcgacattattttatgtgatagatagcatt
449 V T A W G R Y T T I T T A Q K C F D R I I Y C D T D S I
3580 catctagtaggaacagaagttccagaagcaatcgatcacttgggtgatcctaaaaaacttgggtatgggggcatgaaagcaca
477 H L V G T E V P E A I D H L V D P K K L G Y W G H E S T
3664 tttcaacgagcaaaaattcattcggcagaaaacatacgtagaagaattgatggcgaattaaatgtaaagtgtgctgggtatgcca
505 F Q R A K F I R Q K T Y V E E I D G E L N V K C A G M P
3748 gatcgaataaaaagagattgtaacttttgacaatttgaagttgggttttcaagctatggaaagttgctacctaaaaaacacaa
533 D R I K E I V T F D N F E V G F S S Y G K L L P K R T Q
3832 ggtggcgtgggtattagtagacacaatgtttacaatcaataa 3873
561 G G V V L V D T M F T I K *

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11305 atggaagaacgaattgatattcaaatgaacaagatgaagaagaaaatcaaaagaattacatttgcaccctgaaacgaaacccg
1 M E E R I D I Q M N K M K E E N Q K N Y L L H P E T N P
11389 aaacaagttgttttgatgaacattgcattgaaatgaaatcaggagagtttcaacaattttgtgacacagaagaaaatgaca
29 K Q V V P D E T L H G N E N Q E S F N N F V D T R K M T
11473 actacaattgatgtaagtgccttgggggtatcgctgacgggtgaacagattgtacaccaatattaaataaattacttgaagaa
57 T I D V S A Y G V I A D G V T D C T P I L N K L E E
11557 aaaagcgaatgggtatcactttttatcttctccttctgtgaacgtgattcatattatcgctttgctaacaccattgaattgaaa
85 K S E M G I T F Y F P P C E R D S Y Y R F A N T I E L K
11641 cgtgatcactgtatgttactttcttaggatcgggagaaacgacattaaagtttgaacaaatgacggcatttaattgtaaacatc
113 R D V P V V T F L G S G E T T L K F E T M N I
11725 gaaagtttcaatattgatggttttgcattatggttgcacaaaggcgtcaaaagtggtaaaggaattttctttaatgatactgc
141 E S F N I D G F A L W L P Q G A Q S G K G I F F N D T R
11809 aattacaactcgttttgactttgtttgttctgtaacttttaaatgaaggaaacgtatgttggctgtagaggttaga
169 N Y N R F D F D L F V R N C T L N E G T Y V V V A R G R
11893 ggggttacatttgaataattgtctattctctaatatctctcaagcaattatcaaaacagcttttcccgtgtaaatgtatgtgg
197 G V T F E N C L F S N I S Q A I I X T A F P D V N G M W
11977 caaggggaacgatataactaggggtacaggttttagaggtttcttctgtgaaaaaacaacggtattcattttgtacagcgatc
225 Q G N D I N T R G T G F R G F F V K N N R I H F C T A I
12061 attatcgacaatgacgatgattatcagaatgttaatttctgtgaaatttctggttaacacacaatcgaaaggtggcgtaagttat
253 I I D N D D D Y Q N V I N F C E I S G N T I E G G V S Y
12145 tatcgaggatagcgcataacttgcattgccaanaacacattttcttagcatcggaaatagaaacgctttgttfgagttt
281 Y R G Y A H N L H V Q N N H F L A Y G N R N A L F F G
12229 caagatgtggatcaagcttatattgatgtatgtttattgtcgttaactcacaagtcgaggggaatgaatagtagcagctatttca
309 Q D V D Q A Y I D V D V Y C R N S Q V E G M N S T A I S
12313 cgtttaattgttggtaacggacattaccgaaacttaagattacaggttaaatatattcgttgcgaaggacgttatcacggtg
337 R L I V V Y G H Y R N L K I T G K L Y R C G H V I T L
12397 tatggcgggtggcgttaatttctattgtgacttgatggcacaagaagcacctttgacggacgggtaccggtttattcaaacggct
365 Y G G G V N F Y C D L M A Q E A P L T D G Y R F I Q T A
12481 gacaactcaggttaactatgaggttgggttctgctggttctgaactcaacaaagtaaatcaccaatgatctataaagca
393 D N R V N Y D G F V V R G L S N S T K V N T P M I Y K A
12565 cctcagactgttttctataatcgtagaatcgatcgtgtaacaggtccaatgcaagtaatgtatataactag 12639
421 P Q T V F Y N R R I D H V L T G P N A S N V Y N *

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4626 atggctgacaaaatcacagaacaagatgttctcgtgcacaaaatgtagaacaccagtagcaattaatgactgtatttataat
1 M A D K I T E Q D V L R A T N V E T P V Q L T A T A I Y N
4710 agttcatcatctcttttccaggcgaacgtacctatgcaaaatgcagataaacatcgaagcgggtgggtgacgggatcacacgttta
29 S S S S L F Q A N V P M P N A D N I E A V G A G I T R L
4794 gacgtagtaaaaaacgaattttatttcaacttttagttgacgggtattggttaaagtagttatccgatacaaatcttggcgtaacct
57 D V V K N E F I S T L V D R I G K V V I R Y K S W R N P
4878 ttgaaatgtttaaaaaaggaaacatgaccttaggtcgaacgtatgaagaatttttggatgacattgcacaggaacataagttc
85 L K M F K K G N M P L G R T I E E I F V D I A Q E H K F
4962 aacctgacgagtcgtttacaggggtattttaaagggaagttcccgatgtaaaaaacattgttccacgaataatcgtgtaaggt
113 N P D E S V T G V F K Q E V P D V K T L F H E I N R E G
5046 tactacaacaaacgatccaagaagcatgggttagaaaaagcatttacttcatgggataatttcaatagtttctggtggtgta
141 Y Y K Q T I Q E A W L E K A F T S W D N F N S V A G V
5130 atgaacgctttatacacaggtgacgaagtaagcgaatttgaatacacgaaattattaatagcaaaactaccaagaaaaagagcta
169 M N A L Y T G D E V S E F E Y T K L L I A N Y Q E K E L
5214 ttcaaaagagatcgaatttggcgaatttactgaatcaaatgcaaaagatttatccgtaagatcaaatcaactcaacaaatta
197 F K E I E I G E I T E S N A K E F I R K I K S T S N K L
5298 gaatttatgagttccgcttacaacgctcaggaggttaaacatctactcctcaaatctgatcaatagcttatattgacggcgac
225 E F M S S A Y N A Q G V K T S T S K S D Q Y V I I D A D

5382 acagacgcaaccattgacgttgacgttttagcagcggcattcaatatgagtaaaactgactttgtaggacacaaaactcgttatt
253 T D A T I D V L A A A F N M S K T D F V G H K I V I
5466 gatgagtttctaaaaaagggaaggaatcgtaaatattgtggcagttattgtagatagtggaatgggttatgactacgac

320

281 D E F P K K E G E E S S N I V A V I V D S E W F M I Y D
5550 aaattgtacaaaacacagctctatacaaccctgaagggttatattggaattattggttgaccaccaccaactatattctact
309 K L Y K T T S L Y N P E G L Y W N Y W L H H H Q L Y S T
5634 tctcaattcgggaacgctgttctttgttaaatcagcaacaaaacctgtcacaaaagttgcttttgcagtgcaacaactagt
337 S Q F G N A V A F V K S A T K P V T K V A F A S A T T S
5718 gttgttaaggatcatctaaagatatcgcatgtacatttacaccagtagaagcaacaaacaaaggaaggtgttcatca
365 V V K G S S K D I A L T F T P V E A T N Q Q G E V V S S
5802 gcaccagcattggttaaggcaacccgtaaaacaaacagcaggttaaggcactgccgaacagtagaaggcttagaagtcggtcaa
393 A P A L V K A T V K Q T A G K A T A V T V E G L E V G Q
5886 tcattagtaacattcacagctatcggaaggtcaacaagcaacggttctgttacggttacttctgactaa 5954
421 S L V T F T A I G G Q Q A T V L V T V T S D *

182ORF005
12651 atggcaactcttacaaatgaacaaatagctagaggacaacaaatcgctaaaatactttcaaaatatggctataataaaaaattca
1 M A T L T N E Q I A R G Q T I A K I L S K Y G Y N K N S
12735 caagtaggagttgtcgccaatctccattgggaatcggtgttgaacccgaacagcaatgaatatggtggaggcggtatggg
29 Q V G V V A N L H W E S A G L N P N S N E Y G G G G Y G
12819 ttggtcaatggagcgtctaaaagcaatctttatcgcaagcacaaaatttgggtgtgtctaatgctaaagctgaaacgtgtggaa
57 L G Q W T P K S N L Y R Q A Q I C G L S N A K A E T L E
12903 ggtcaagcagagatcatcgctcaagggtataaaacaggtcaatggatgataatacacctgttcttctcgaggttataactaac
85 G A E I I A Q G D K T G Q W M D N T P V S S A G Y T N
12987 cctcagaccctttcagcatttaacaaatctgcaaatattgatgtgtctacaattaattttatgtgtcactgggaacgcctcggt
113 P Q T L S A F K Q S A N I D V A T I N F M C H W E R P G
13071 aaacttcataatgaagaagactttgatcttgcaagcttatagtaagcatattgacggtagcgggtggcggtggcgtaaacgt
141 K L H I E E R L D L A Q A Y S K H I D G S G G G V K R
13155 tgctatggaaaccccaatcaagaatacaaatcttgatcctaaaagtttcatgagtggaacaacttttggcagcgtgacggaac
169 C Y G T P I K N T N L D P K S F M S G Q L F G T H A G N
13239 ggcagaccaataatttccatgatggtttggacttttcaattgatcacccctggcaatgaaatgattgcattgttcgatgga
197 G R P N N F H D G L D F G S I D H P G N E M I A C C D G
13323 acagtaacacatggttgaacaatgggagcattaaagcgtattttgtgataaatgatggtacttacaatatcgtttatcaagaa
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13407 tttagtataaccagtcataataaaggtaaaagttggcgacaaaagttgaacaggaacagtttgcgcaatcgtgacgaggat
253 F S Y N Q S N I K V K V G D K V K N G Q V C A I R D A D
13491 cattacatttaggttttactaaaaagattttatcgactcggttaggatcttcttctcatagatgatggaacatgggaagaccct
281 H L H L G F T K K D F M T A L G S S F I D D G T W E D P
13575 ttgaagtttttagggcaatgttttgagatggagatctggcgagataatgacgataacaataaggataaaaatgatcttatt
309 L K F L G Q C F G D G D T G G D N D D N N K D K N D L I
13659 tatctattgctatccgatgccttgaatgggttgaaattttaa 13700
337 Y L L L S D A L N G W K F *

182ORF006
14995 atgacaaatagcttaggcgttaaaacttgaagagaaaaacttataactataaccctaacaatgcttttaggttttaattgcctaagt
1 M T N S L G V K L E E K N L Y Y N P N N A L G F N C L M
15079 ttggttgaataggcgacgtggtataggttaaaacttattggttataaaaaatttgttgaatcgctttattaaacacggcgaa
29 L F V I G A R G I G K T Y G Y K K F V V N R F I K H G E
15163 caatttatttatttaagaagattcaaacagaacttaaaaagattcctcaattttcaaaaacaaatggcgaaagaatttctgat
57 Q I Y L R R F K T E L K K I P Q F F K T M A K E F P D
15247 cataaacttgaagtaaaaggaaaagaattctattgtgatgataaataatgggttgggctgttccacttagctagctgggaatt
85 H K L E V K G K E F Y C D D K L M G W A V P L S T W G I
15331 gaaaaatctaataatccccgaagttcgtacaattttgttgatgagtttttaattgagaaatcaaaaactcattttacca
113 E K S N E Y P E V R T I L F D E F L I E K S K I T Y L P
15415 aacgaagctgaagccttattgaacatgatggaacgggtttccgaagacgtacaaatacaagatgtgttatgttgagtaatgca
141 N E A E A L L N M M E T V F R R R T N T R C V M L S N A
15499 actagttagtgaaaccttatttctgtatttcaatctgcagccagatttgaataagcgttttcaatcatcaagatcgaggt
169 T S V V N P Y F L Y F N L Q P D L N K R F N L Y Q D R G
15583 atattgattgaattgtgtgattcaaaagactttgcagaagtgaaagagagaaacaccttttggtagattgattcggtggaacagaa
197 I L I E L C D S K D F A E V K R E T P F G R L I R G T E
15667 tacgaagatttttagtatcaacaatgagtttgcataatgatagtgatagctttattgaaaagagaagtaaaaatagtagtttctta
225 Y E D F S I N N E F V N D S D T F I E K R S K N S S F L
15751 tgcgccattgcttttgaagggaatactttgggtattggatagacgctgaaacaggttgtgtctatgtgagttatgattatcaa
253 C A I A F E G K I F G Y W I D A E T G C V Y V S Q
15835 ccaatacaaatcatttttatgcaatgactacgaagaccatgaagaaaatagattgctgatgaaaaattggcgaaataattat
281 P N T N H F Y A M T T K D H E E N R L L M K N W R N N Y
15919 tatctttcaacagtgggcgaaagcattcaagaatagttatctcggttttgataacattgtttattagaatttacattatgattg
309 Y L S T V A K A F K N S Y L R F D N I V I K N L H Y D L
16003 ttttaaatagatgaaaaatctggtaa 16026
337 F N K M K I W *

182ORF007

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1 M S R R K G A G L A R N N Y T A K S R P Y P N F P Y S
7879 agtgatgtagaagaatcagctactatgaacattatcgtagacaactcacgctccttacgtttggttggttggaataat
29 S D V E E I S Y Y E H Y R R Q L T L L T F Q L F E W E N
7963 ttgcaaaatcaattgacctcggttatttagaaattgctttacacactaatgggttatcttgggttcttttaagacctacact
57 L P K S I D P R Y L E I A L H T N G Y L G F L D P T L
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321

85 G F M V C A G A E D G Q I D H Y H N P I F F T A N E A M
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141 V P T L P S L H R F A L D M A D I N Q I S R V N R R A Q
8299 aaaacacctgtaattattcaaacgtgatgaaagaataacttctcattgtctacaagcttataaccaaatgacgaaaaataatcag
169 K T P V I I Q T D E K K Y F S L L Q A Y N Q I D E N N Q
8383 gctgtttttgtggataaagatatggagtttgacgaatcttttaattgtatggcaacaatgctccatatgtagtagataaacta
197 A V F V D K D M E F D E S F N V W Q T N A P Y V V D K L
8467 cgatcagaattgaacgaagtattggaatgaagtgttaacttttctaggtatcaacaatgctaactgtagataagactgcacgtgta
225 R S E L N E V W N E V L T F L G I N N A N V D K T A R V
8551 caaacatcagaagttcttatcaacaatgaacagatgaaagtctcaggtaacatctgtttaaatacaagaaagagttttgcat
253 Q T S E V L S N N E Q I E S S G N I L L K S R K E F C D
8635 cgtgtaaatcgtgtctttggcgatgaacttgacggaaagattgacgtgaagttagaacagacgcgttcgacaattacaactg
281 R V N R V F G D E L D G K I D V K F R T D A V R Q L Q L
8719 gggcgaggtcaatcaaaaaagaccagatgagtgagggttgccaagtgtactactaa 8775
309 A A G Q S K K D Q M S G G L P S A T *

182ORF008
14105 atgatgaatggattatctctagttatcaaacggaattgatctttcaaaagttccatgcatgtttgttaaatattaaagca
1 M M N G I D I S S Y Q T G I D L S K V P C D F V N I K A
14189 acaggcggaacaggttatgtaaacctgattgtgaccgagcatttcaacaagctttgtcttttagttaaagattgggtgtgat
29 T G G T G Y V N P D C D R A F Q Q A L S L G K K I G V Y
14273 cattttgcatgagaggggtttagaaggtacacctcaacaagcgcaattcttttagataataatgaaggttacattgggt
57 H F A H E R G L E G T P Q Q E A Q F F L D N I K G Y I G
14357 aaagctgttcttattctgactttgaaggtcaaatcagaagatgtaaatggcgaaagcatttcttgattatgtttataat
85 K A V L I L D F E G S N Q K D V N W A K A F L D Y V Y N
14441 aaaacaggcggttaagcatgggtttatctgtatcacgcaaacctcaatacaactgattttctagtattgcaaaagcgattat
113 K T G V K A W F Y T Y T A N L N T T D F S S I A K G D Y
14525 ggtttatgggttgctgaatatggatcaaatcaaccacagggctactctcaaccagcgccacctaaaaaataattttccaatt
141 G L W V A E Y G S N Q P G Y S Q P A P P K T N N P P I
14609 gttgctgttttcagtttacaagtaaggacgtttaccaggatacaacggcaatcttgatttgaatgtttctatggcgatgggt
169 V A C F Q F T S K G R L P G Y N G N L D L N V F Y G D G
14693 aatcacatgggatctgtatgtaggtaaaaaacaggtataaatgttctctcgtgaaaataaataattttgacgccacaagtgatgag
197 N T W D D L Y V G K K Q D Q I V P P E N K I F D A T S D E
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225 F I F T L T T G S T S V F Y F D G E T I F E L S K D P T C D
14861 ctcgatcatattagaggaacatacatcatgttcatggaaaagaatcccatcaatgggtggacacctgaacaatttgatatt
253 L D H I R G T Y N H V H G K E I P S M V W T P E Q F D I
14945 tacttaaaatgtatgaaaagaaccagtatataatag 14983
281 Y L K M Y E K K P V Y K *

182ORF009
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1 V L L K R Y I E S P T Y Y Q P E L S R K E R I E V G R K
8849 caattgtttgattttgattatcgttttatgacgaacaaacagcagcaatttgaaacaaaatttatcaactcttttacttg
29 Q L F D F D Y P F Y D E T K R A E F E T K F I N H F Y L
8933 agagagatggtcagaaacgatgggattcaattcaattcttgacgaatattttaaacttaaacatgcccatttggaataaaa
57 R E I G S E T M G S F K F N L D E Y L N L N M P Y W N K
9017 atgttcttatcaaatcttgaaagatttccgatttttgatgacatggactacaccattgatgagaaacagaaattgttaaatgag
85 M F L S N L E E F P I F D D M D Y T I D E K Q K L N E
9101 attgatacaaacatcaaacggaatcgtgatgaatcgaagaacaaacgaagcaagtagatcaaacagacacagaaacaaaaat
113 I D T N I K A N R D E S K N Q T K Q V D Q T D N R N K N
9185 acacgtgacacaggaacacagatttcttcaaggaacactttatcacagacacctcaaaaagatttgagaattggcagcaat
141 T R D T G T T D S F S R N T Y T D T P Q K D L R I A S N
9269 ggagatggaacaggtgtaataatcgaacaaatatcacagaagatttgagtaagaacaaacaggtccacagggcggtgaa
169 G D G T G V I N Y A T N I T E D L S K E T T S T G V E
9353 acaaaacagcaaaaacaaatcaaaatcacgaagcaatgcttctgaaaaagaaacaaagaacacagacatttaataaagatcaa
197 T N N D K T N Q N T R S N A S E K E T K N T D I N K D Q
9437 aatcaaaccaagatcagattacacgatataaaggtaaaaggggaacactgattatgctgacttactcgaaaaattcgtaga
225 N Q T K D T I T R Y K G K A K G N T D Y A D L E K Y R R
9521 agtgttttgagaattgagaaaaatgatcttttagaagaatgaacaaggaaggcttatttctccttgtttatggaggggagtag
9601
253 S V L R I E K M I F R E M N K E G L F L L V Y G G R *

182ORF010
1310 ttgaccgtaagaatatcaagaatgatagagccaagttagagaaaatctacggttaaatctaacaaagctcgtaaaaaatacaat
1 L T V R I S K N D R A K L E K I Y G K S N K A R K K Y N
1394 cgttttaagacaaaaggagttgaggaaggcaacttccaactgttccaacatcaagaaaagacttactgactcgaataatca
29 R L R Q K G V E E R Q L P T V P T S K K R L I D Y V K S
1478 acaaatagagtcgtatgtatttcaacaagattgtagacaggttggttagattttgcacacaccttacaacgagaatttaattttt
57 T A N M S R S D F N K M L D E L V D F A Q Y P Y I F
1562 gagatcaacaagcgaatgttgcaatctcaagagcgcaaatcaagaagcgcaaatcaaacagagcaagctcaaaaagcgaaa
85 E I N K R N V A I S R A Q I K E A Q I K T E Q A Q K A K
1646 gaagaacactacaagagcttaacaaagttgaagttgaagagccacagaaaaacacaaattgtcacacctatttttaacagag
113 E A H Y K E L N K V E V K K P T E N T I V T P T I L T E
1730 ttagggtgctgacttaccttttcaagcaataccagattttaaatttgacgctttcacttctccagaaggagttcagttctattta

322

141 L G A D L P F Q A I P D F N I D A F T S P E G V Q S Y L
1814 gaaaatataggaaaacagacgaacaatattttgacgaaagagaccaactttattacgacaatttcagacaagcgatgtttact
169 E N I G K Q D E Q Y F D E R D Q L Y Y D N F R Q A M F T
1898 attttcaattcagacgctgacgatattgttcgtttacttgactcaatggggcttgatctatttatgaaaacatatgttagtaac
197 I F N S D A D D I V R L L D S M G L D L F M K T Y V S N
1982 ttcttagacatgaaccttgactacatttatgacgaagcagaagtacaacagaaaaaagaacaagtttacagtaagattgcaaaa
225 F L D M N L D Y I Y D E A E V Q Q K K E Q V Y S K I A K
2066 gtgatcgagtcgtaaacaggtggagaagtccctcatataaccccacgaagaacatcacaattaattcagaaacaggagaagaa
253 V I E S E T G G E V P S Y N P T K N I T I N S E T G E E
2150 ttatga 2155
281 L *

182ORF011
9607 atggtagattttaaccccgacaagcggtttgacggtttacccgctgtattcaaagaacgcttttagcaaatatcctcactatgaa
1 M V D F N P D K R F D G L P A V F K E R F S K Y P H T E
9691 tacagatatgaattactattagatgaagaagtatcggttttaattgcctatctgaatgaagttgggtgcttttagttaatgatag
29 Y R Y E L L L D E E V S A L I A Y L N E V G A L V N D M
9775 agtggcttatttaattactttatcgaaacattttgttgagaagttagaagatcacaaatgacacactcaaaaaatgggtgtct
57 S G Y L N Y F I E H F V E K L E E I T N D T L K K W L S
9859 gatggtagcttagaaaaatttaataatgatactgttttgcaaatatatcaaaagaatcaaaagattacaaatccttggttgct
85 D G T L E N L I N D T V F A N Y I K E I K R L Q I L V A
9943 gaaacacgtgctaacagtgatgaatattcttttgcaaaaaataaacggatgttgctgatgcgaacattttggtataagatt
113 E T R A N S V N I L L T K N K P D V A D D R T F W Y K I
10027 caacgcgacaataactgattatggagccgactctattgacacgcttacgtattgttgcaatcaataaagtttagtgggtggaatacc
141 Q R D N T D Y G A D P I D T L R I V A I N K V S G W N T
10111 gctacaggagatatttatcttaacattaaaggaacggagggtgtataa 10158
169 A T G D I Y L N I K G T E G V *

182ORF012
10872 atggcaataaaaaatattcaaatgaaggatagcaatgacaataatttatatccaagtgttcgagcagaaaaacttgtagatttg
1 M A N K N I Q M K D S N D N N L Y P S V R A E N L L D L
10956 accagtcgtgctgaattaacaatgacaaattgtcaattatgacgctggtgataaaacaaatgcaattcttctcggtgca
29 T S R A E L T M T N C Q L Y A A G D K T N A I S Y L G A
11040 gtaggtagctcgaaggatgataaagtgttactgaaagtttgacaaacctgtgatcacacgctaccagaaggttttagacca
57 V G M L E G M I K F T E S L T N P V I T T L P E G F R P
11124 ataagaacaaaacgtattggtgtttcgcaaaatattacacaccaaattccaacagatacaaaagaagtgtttatgtatcaatc
85 I R T K R I G C F A K Y Y T P N P T D T K E M V Y V S I
11208 acacctgatggcaagtaactgtaaatgacaatgtaggtaaaatcgaatatctatccctagataaattgcgttttcctctaaaa
113 T P D G K V T V N D N V G K I E Y L S L D N C V F P L K
11292 taa 11294
141 *

182ORF013
10456 atggcagataaaaaatattcaaatgcaggataaagatcataatcggttaattgctgtttacaattgctaaaaatggttctaacaggc
1 M A D K N I Q M Q D K D H N R L M P V T I A K N V L T G
10540 gactctaactctgaattagttaattgctgaaataagaggtaacgctagtgaagctaaaacacttgcaacaacagctaaagaaact
29 D S N L E L V N A E I R G N A S E A K T L A Q Q A K E T
10624 gctgctggtttgtcaacagaaattgacacagtaaatcaacgcgaaatcaagcgttgacgaaggtggtacagcacaacaaacc
57 A A G L S T E I D T V T S T A N Q A L T K A G T A Q Q T
10708 gcagaacaagcgaacaaacagcaaacagtatcagcgcagttgcaacggcagctaaaaacacagctgattcagcacaacaaagt
85 A E Q A K T T A N S I S A V A T A A K N T A D S A Q K S
10792 gcaactgatctagctggttcgagtaagcagtttagaggacacagcaatacataactgtattaccatag 10860
113 A T D L A V R V S S L E D T A I Q Y T V L P *

182ORF014
13716 atgatagaatatatcacacaatgggtggcagatgataatcatctgtttatggtttgattatatggttaattggttgcaatgatt
1 M I E Y I T Q W L A D D N H L V Y G L I I W L M V A M I
13800 atcgattttgtggttaggttttacaattgcaaaatttaacaaggaaatcgacttagtagtttttaagctaaagcaggtatcatt
29 I D F V L G F T I A K F N K E I D F S S F K A K A G I I
13884 gttaaggtggcagaaatggttttagtgggttactttattcctgtagcagtaaaatcggtgcagtaggtattacaatgtatata
57 V K V A E M V L V V Y F I P V A V K F G A V G I T M Y I
13968 acaatggtggtttggtttgattttatcagaaatttatagtatactaggacatatctcagatatcgatgatgataaattggact
85 T M V G L I L S E I Y S I L G H I S D I D D D N N W T
14052 gattatgttaagaagtttttagcggaaactcaacagaaaggacgatattaatga 14108
113 D Y V K K F L D G T L N R K D D I K *

182ORF015
854 atggaaatcgtaaaaagcacatttgacacacaaacaccagaaggaattgtacaagtattcaatgccacaaacggggcttcaatt
1 M E I V K S T F D T Q T P E G M L Q V F N A T N G A S I
938 ccgttacgtaacgcaattggcgaagtactagaaatatttagttactcagacgaagtttctggtttggtggagcc
29 P L R N A I G E V L E L K D I L V Y S D E V S G F G G A
1022 gaaccatcacaaagcagaacttagtcgctttctcacagaagatggttaaaccttatcggtgtgatcagcagtagcaacaaatca
57 E P S Q A E L V A F F T E D G K T Y A G V S A V A T K S
1106 gctaaaaacctaattgatgatgactgtaacccctgacatcaaaccaaaatttctttgtcggaaggaaatcaaacgggtgga
85 A K N L I D M M T A N P D I K P K I S F V E G -K S -N- -G
1190 caaaaattgttaattctacaagtggtttcactgtag 1225
113 Q K F V N L Q V V S L *

182ORF016
17033 atgattaacaatttatcattaatttttaggggttttaactaaactaaagatgacaacgatagtttagcgtctatcaagtca
1 M I N N L S L I L E G G L N Q L T K D D N D S L A S I K S
16949 gaaataacacaaggaggaaacaatttaattttatcattgattacgtttacaaaagagttcgtgttaacacatgataaataaac

323

29 E I T Q G G K Q L I L Y I D Y V T K E F V L T H D K Y N
16865 tatgtttatcttgatagccattgcattaatatcgcaataacgaaatcaatgaaaagcgttgaaactactatcggaacaattgaaa
57 Y V Y L D S H C I N I A I T K S M K S V E H Y A E Q L K
16781 catgacggatataaacaattacggacaaatag 16749
85 H D G Y K Q I T D K *
182ORF017
154 atgaaatattcactacaacaatagatgaaattaaatcaacaattttcagaattagattaaaaaggcatgaactagaggaattg
1 M K Y S L Q Q I D E I K S T I P R I R L K R H E L E E L
238 gtggacgaagtaaacgatattgctaaagatccggaggaaagatatctttatcgttttattacacagaagaagaacgtttgtt
29 V D E V N D I A K D P E E R Y L L S F Y Y T E E E R L F
322 gaaattccctctgcaagattaatagattattacaacgaaaagatcacaaatctgaaatcggaatcatatcactcgaaaaaga
57 E I P S A R L I D Y Y N E K I T N L K S E I I S L E K R
406 ttacaaaaactagtaaaataa 426
85 L Q K L V K *
182ORF018
16737 atgattgcacgaacattcaagaacaccgcgaactaattgaatgggttacgtttctactgtaaacgtaacctttcagacaatgaa
1 M I A R T P K E H R E L I E W L R F Y C K R N L S D N E
16653 aaaatagagcatagaggggactttacaagatttcgacgttccggaataaatatcacgaaactttgttaactcattcaacg
29 K I E I I E G T L Q D F D V P E I N I T E L L L T H S T
16569 ctattaccggaatcgagtcaatttaacattcttgaaaagtattgtcaggcaatgaaattagtaacttcatacgtaaaagttgg
57 L L P E S S Q F N I L E K Y C Q A M K L V T S Y V K V G
16485 tctcgctatcagtttagcggtacaaatacgaagggtctatttaaggaggtggaataa 16429
85 S R Y Q L A L Q I P K G Y L K E V E *
182ORF019
4323 atggaaattaaagaacatgaatcaatttttaaatgggtattcttgaaagtgtcacagacgggtgaagcaagatcaaagattgtagaa
1 M E I K E H E S I L N G I L E S V T D G E A R S K I V E
4407 catcttgagcattgagagaagactacggagcaacaactgaagctttgacatcagcaaatagcacacttgaaaagttaaagaaa
29 H L E A L R E D Y G A T T E A L T S A N S T L E K L K K
4491 gataacgaagcgttggtatttcaaaactcaaaattgttccgagaacgagcgatcgtagaaccagcagaaaaataacgaaccagaa
57 D N E A L V I S N S K L F R E R A I V E P A E N N E P E
4575 acagaccagaatattacactagacgatttaggaatttaa 4613
85 T D Q N I T L D D L G I *
182ORF020
10158 atggcagacattagaacacaactaacaagtgaagatggatcagacaattttttccaatttcaaaagccgttaattattatgact
1 M A D I R T Q L T S E D G S D N L F P I S K A V N I M T
10242 aatagcggtagcaatgtagaaggagaattgggtacactcaaaacaaatgacgaaacaatgaatacctcagttcaaaatgctgt
29 N S G T N V E G E L G T L K Q N D E T M N T S V Q N A V
10326 gttactgccaatcaagcaaaagattctgtagctgaattaaatgtaaatgttggttaaactaaccaatcgaataacaacattagag
57 V T A N Q A K D S V A E L N V N V G K L T N R I T T L E
10410 agtacagtggttaattcttgatgggtattcgttagaggtgtaa 10454
85 S T V A N L D G I R Y V E V *
182ORF021
17339 atgaacaataaatcattaatagctgaaaaaggagaggtatctctacttcacccctttaatgagtgaggatgaattatcatatc
1 M N N K S L I A E K G E V S L L H P F N E W D M N Y H I
17255 atagataccgaaaaacaataaactattatcttattgataggttaggcgatgaggaatattgtttgttatcttttgaagaa
29 I D T E N N K H Y L I D I D E V G D E E Y C L L S F E E
17171 ctaaaaggaattagatattgattcttattccgagttattcgtgaaaactacagaaataacatatata 17106
57 L K E L D M D L I S E Y S W K T T E I T Y *
182ORF022
12868 gtgggtgtgtctaatgctaaagctgaaacgttggaaggtcaagcagagatcatcgctcaaggggataaaacaggtcaatggatgg
1 V G C L M L K L K R W K V K Q R S S L K G I K Q V N G W
12952 ataatacacctgtttctctgcaggttataactaacctcagacccttcagcatttaacaatctgcaaatattgatgtgtgcta
29 I I H L F L L Q V I L T L R P F Q H L N N L Q I L M L L
13036 caattaattttatgtgtcactgggaacgccctgggttaaacttcatatcgaagaaagacttgatcttcgacaaagcttatagtaagc
57 Q L I L C V T G N A L V N F I S K K D L I L H K L I V S
13120 atattgacggttagcggtggcggtgta 13149
85 I L T V A V A V A *
182ORF023
12189 atgggtgtgtgttttgacatgcaagttatgcgcataatcctcgataataaacttacgccaccttcgattgtgttaccagaaatttc
1 M V V V L D M Q V M R I S S I I T Y A T F D C V T R N F
12105 acagaaatttaattacattctgataatcatcgctcattgtcgataatgatcgctgtacaaaatgaatacgggtgtttttcacaaa
29 T E I N Y I L I I I V I V D N D R C T K M N T V V F H K
12021 gaaacctctaaaacctgtaccctagattgatatcggtcccttgccacatacattttacatcggggaaaagctgttttgataat
57 E T S K T C T P S I D I V P L P H T I Y I G K S C F D N
11937 tgcttgagagatattagagaatag 11914
85 C L R D I R E *
182ORF024
6174 atgcttgtaactatctcatcttttaaaacgaagaacttatcctagtaaatggcagtagtgcctttgttactgatattgaataa
1 M L V T I S S L K T K K L I L V N G S M P L L L I L N I
6258 agaataacacacaagtttcgttacctttgaaattgatgttttacaaacttatcgtttcgatattggtataggaagatttca
29 R M T T Q V S L P L K L M F Y K L I V S I L V Y E K V S
6342 ttgcaaaagaacacctcaactttattattcgaatggaatacctttcattaatacaattgaagagtcgcttgattacggttagag
57 L Q K N T L N F I I R M E Y L S L I Q L K S R L I T V E
6426 aatacacacaacaatgtaa 6446
85 N T Q Q Q M *

324

182ORF025
548 atgggtcgaaaactaatgcaacgaaacgtaacatcaactaaagtagaattctcagaagttatcgtaacagatggagcgccaaca
1 M G R K L M Q R N V T S T K V E F S E V I V Q D G A P T
632 attgtaccatgcgaaccagttgtcttaacaggaactttcagaagaaaagctttatcagcgatcaaacgtaaaaacctgat
29 I V P C E P V V L T G K L S E E K A L S A I K R K N P D
716 aaaaacgtagttgtaacaaatgtttcacatgaaacagcgctttacacaatgccagtcgataaatttatcgagtttagcagacaaa
57 K N V V V T N V S H E T A L Y T M P V D K F I E L A D K
800 tcaacacaagcctaa 814
85 S T Q A *

182ORF026
13259 atggaaattatttggctctgccgtttctcgtcatgcgtgccccaaagtgtccactcatgaaacttttaggatcaagatttgtatt
1 M E I I W S A V S C M R A K K L S T H E T F R I K I C I
13175 cttgattggggttccatagcaacgttttacgccaccgcccagctaccgtcaatgatgcttactataagcttggcgaagatcaag
29 L D W G S I A T F Y A T A T A T V N M L T I S L C K I K
13091 cttttcttcgatataagtttaccagggcgtttccagtgacacataaaaatttaattgtagcaacatcaatttgcagattgttt
57 S F F D M K F T R A P P V T H K I N C S N I N I C R L F
13007 aaatgctga 12999
85 K C *

182ORF027
14896 atgaacatgattgtatgttctcctaataatgatcgagttgtgttggatcagacaattcaaagatcgtttctccgtcaaaataaaa
1 M N M I V C S S N M I E L C W I R Q F K D R F S V K I K
14812 cagcctgtgtctacctgttgaagagtgaataaaactcatcactgttggcgctcaaatatttttatttcaggaggaacaaattg
29 H A C A T C C K S E N K L I T C G V K Y F I F R R N N L
14728 atcctgttttttacctacatacagatcccagtattaccatcgccatagaaaacattcaaataagattgcccgttgcattcctgg
57 I L F F T Y I Q I P C I T I A I E N I Q I K I A V V S W
14644 taa 14642
85 *

182ORF028
14430 atgtttataataaaacagcggttaaagcatggttttatacgtatatacagcaaacctcaatacaactgatttttctagtattgcaa
1 M F I I K Q A L K H G F I R I Q Q T S I Q L I F L V L Q
14514 aaggcgattatggtttatgggttgcgtgaatattggaatcaaccacaaggctactctcaaccagcgccacctaaacaaata
29 K A I M V Y G L L N M D Q I N H K A T L N Q R H L K Q I
14598 attttccaattgttgcctgttttcagtttacaagtaaggacgtttaccaggatacaacggcaatcttgatttga 14672
57 I F Q L L P V F S L Q V K D V Y Q D T T A I L I *

182ORF029
17606 atgaatgaaccgatcgatatacagaaattttatcaataacgtggtatgtatgaaaatttttagagatgaggataaaacttagt
1 M N E P I V Y T E I Y S N N V V C M K I F R D E D K L S
17522 aaattctcttattagaatttgaggtggatgaggctaaaagttacttgaaaataaaacaatttcttgatgataactggact
29 K F L Y L E F E V D E A K K L L E N K T I S F D D N W T
17438 ttctcaataaattatccagaatattaa 17412
57 F S I N Y P E Y *

182ORF030
16429 atggctacattctacaaggaaccaatatacagatatacacagtattttatatagatggttgggaggttttgatacacaacccgaa
1 M A T F Y K E P I Y D I T V F Y I D G W E V L I H K T E
16345 cctctcaccttaacaaaagcatttaaaatagccgtatatacctagaatggatatagtgaattgctgttagaataagaaat
29 P L T L T K A L K Y S R I Y L E M D I V N C V R I E R N
16261 ggagcgtcctatagctacattttacaggggaattattaaaactgtataaggagaagaactatga 16199
57 G R P I A T F Y R E L L K L Y K E K E L *

182ORF031
8603 atgttacctgaactttcaactgttctcattgttagataagacttctgatgttgtacacgtgcagttcttatctacgttagcattg
1 M L P E L S I C S L L D K T S D V C T R A V L S T L A L
8519 tgcatacctagaaaagttaacacttcattccatactcgttcaattctgatcgtagtttatctactacatagagcatttgg
29 L I P R K V N T S F H T S F N S D R S L S T T Y G A F V
8435 tgccatacattaaaagattcgtcaactccatattctttatcccaaaaaacagcctga 8379
57 C H T L K D S S N S I S L S T K T A *

182ORF032
11413 atgtttcatcaaaaacacttgtttcggttgcgtttcaggggtgcaataggttaattcttttgattttcttctttcatcttgttca
1 M F H Q K Q L V S G S F Q G A I G N S F D F L L S S C S
11329 tttgaatatcaattcgtttctccatataaacctccttatttttagagggaaaacgcaattatctaggatagatattcgatttta
29 F E Y Q F V L P Y E P P Y F R G K T Q L S R D R Y S I L
11245 cctacattgtcatttacagttactttgccatcaggtgtgattgatacataa 11195
57 P T L S F T V T L P S G V I D T *

182ORF033
4942 atgtcaacaaaaattttcttcaactcgttcgacctaaaggcatgtttccttttttaaacattttcaaagggttacgccaagattg
1 M S T K I S S I V R P K G M F P F L N I F K G L R Q D L
4858 tatcgataactactttaccaatacgggtcaactaaagtgaataaattcgtttttactacgtctaaacgtgtgatccctgca
29 Y R I T T L P I R S T K V E I N S P F T T S K R V I P A
4774 ccaaccgcttcgatgttatctgcatttggcataggtacgttcgcctga 4727
57 P T A S M L S A F G I G T F A *

182ORF034
6160 gtgtttatctactctaaaaactccccaggttgtgtatcccttttgataagaacaattcttattctcgttaagaacaggaaacga
1 V F I Y S K N S P E L C I P L I R T I S I L V K N R K R
6076 attaaagtacattcctgttctcgttgaattttaaaccatcttgtgtgtataggtgttatcaaaaggcagcttagccaacaa
29 I K V R F L F L L S F K P S C V C I G V I K R H V S Q Q
5992 ttttacatttgtataccttcttgcataattgtcctccttag 5951

325

57 F Y I C I P S C H N C P P *
1820RF035
15758 atggcgcatagaactactatcttttacttctcttttcaataaacgtatcactatcattgacaaactcattgttgatactaaaa
1 M A H K K L L F L L F S I N V S L S L T N S L L I L K
15674 tcttcgtattctgttccacgaatcaatctacaaaagggtgttctctcttcacttctgcaaagctctttgaaacacacaattca
29 S S Y S V P R I N L P K G V S L F T S A K S F E S H N S
15590 atcaatatacctcgatcttga 15570
57 I N I P R S *
1820RF036
2315 atgtctgtgctgccttgcatctttacaccactcaaaaaagaatcgatttctaaccgaacgtcatattgtcaacgttgctctata
1 M S V L P C I L H H S K K E S I S K P N V I L S T L S I
2231 tcgcatacgccccacgaccatacacgacaatcggttgagatcagttgtgtttcaaagtcgacgtatatttcttaatacataatt
29 S H T P H D H T R Q S L R S V V V S K S P V Y F L I I I
2147 cttctctgtttctgaattaa 2127
57 L L L F L N *
1820RF037
12280 gtgagttacgacaataaacatctacatcaatataagcttgatccacatcttgaaactcaacaaagcgtttctatttccgtatg
1 V S Y D N K H L H Q Y K L D P H L E T Q T K R F Y F R M
12196 ctgaaaaatggtgtgtgtttggacatgcaagttatgcgcatatcctcgataataacttacgccacacctcgattgtgttaccag
29 L E N G C C F G H A S Y A H I L D N N L R H L R L C Y Q
12112 aaatttcacagaaattaa 12095
57 K F H R N *
1820RF038
14769 gtgatgagtttattttcactcttacaacaggttagcacaagcgtgttttattttgacggagaaacgatctttgaattgtctgatc
1 V M S L F S L L Q Q V A Q A C F I L T E K R S L N C L I
14853 caacacaactcgatcatattaggaacatacaatcatgttcatggaaaagaatcccatcaatgggtgtgacacctgaacaat
29 Q H N S I I L E E H T I M F M E K K S H Q W C G H L N N
14937 ttgatatttacttaa 14951
57 L I F T *
1820RF039
9992 atgttgctgatgatcgaaacttttggtataagattcaacgcgacaatactgattatggagccgatcctattgacacggttacgta
1 M L L M I E H F G I R F N A T I L I M E P I L L T R Y V
10076 ttgttgcaatcaataaagtttagtggtggaataccgctacaggagatatttattcttaacattaaaggaacggaggggtgataat
29 L L Q S I K L V A G I P L Q E I F I L T L K E R R V Y N
10160 ggcagacattag 10171
57 G R H *
1820RF040
16202 atgagaaaaagatttctgtctacattaacacacccgatccaaaagcaaaaaaggcgttagcaaaaatcactaacgccaaagaa
1 M R K D F V Y I N T P D P K A N K K A L A K I T N A K E
16118 ccaaaacaaaactatcgagactacaattactatgttatctactattcatcattgtaataagaactaatcggtgtagcttacta
29 P K Q N Y R R L Q L L C Y L L F I I V I E L I V V A L L
16034 aaatag 16029
57 K *
1820RF041
3886 atggaaactatataaagcaatgtttatcgtagctgaaggtactattgacggttacgatactgaacactatgtagatatttct
1 M E L Y K A M F I V R D E G T I D G Y D T E H Y V D I S
3970 ttacatgactttgaagaaatatatggaaaagaacacgtgaaattgaaagcagtaacattagtaaaaacaggaaatttaaaaaaa
29 L H D F E E I Y G K E T R E I E A V T L V K T G N L K K
4054 taa 4056
57 *
1820RF042
10832 gtgtcctctaaaactgcttactegaacagctagatcagttgctactttttgtgctgaatcagctgtgttttagctgcggttgca
1 V S S K L L T R T A R S V A L F C A E S A V F L A A V A
10748 actgcgctgatactgtttgctgtgttttcgctgtttctgcggtgtgtgtgctgtaccagccttcgtcaacgcttga 10671
29 T A L I L F A V V F A C S A V C C A V P A F V N A *
1820RF043
10652 gtgtcaatttctgttgacaaaccagcagcagtttcttttagctgtgtgtgcaagtgttttagcttcactagcgttacctcttatt
1 V S I S V D K P A A V S L A C C A S V L A S L A L P L I
10568 tcagcattaactaattcaagattagagtcgcgtgttagaacatttttagcaattgtaacaggcattaaacgattatga 10491
29 S A L T N S R L E S P V R T F L A I V T G I K R L *
1820RF044
6457 atgaaaagttgttacattgtgtgtgtattctctaccgtaatacaagcgactcttcaattgtattaatgaaaggtattccatt
1 M K S C Y I C C C V F S T V I K R L F N C I N E Y S I
6373 cgaataataaagttgaggggtgttcttttgcattgaaactttctcgataccaatcgaacgataagttgtga 6299
29 R I I K L R V F F C N E T F S Y T N I E T I S L *
1820RF045
6729 atgaatgggtatacctgtatacgcgttacatacatcccactatcttatttaaaaaagggttcttctgttgtaagaaacgccatg
1 M N G I P V Y D V T Y I P T I L F K K G S F V - N R - N M
6645 tactctccaaattagcattgcctgccccattgtgtgtgttatacctccccacttgattgataggaagtaataa 6571
29 Y S P K L A L P A P F G L Y T S P L E L I G S K *
1820RF046
2372 atgggtttcaaatggtgtaaaagaagcaaaagaagatcgaaacattctccactcatatcaaatatgggtgcaatggtgtgttgg
1 M V S N G V K K Q K I E H S P H S Y Q I W V N G M L W
2456 aaattgtgtgggaagtttaattacacaacacaaaatcaggttaaaacgaaaaagagaatctcgaacaataa 2527
29 K F V G K L I T Q Q Q N Q V K R K K R N L E Q *

326

182ORF047
13353 atgctcccattgttccaacatgtgttactgttccatcgcaacatgcaatcatttcattgccagggtgatcaattgaaccaaagt
1 M L P L F Q H V L L F H R N M Q S F H C Q G D Q L N Q S
13269 ccaaacatcatggaattatttggctgctgctgttcctgcatgctgccaataaagtgtccactcatga 13201
29 P N H H G N Y L V C R F L H A C Q K V V H S *
182ORF048
3395 atgtcagggtttgttccgaactttccatacaagctattttaacatacctttggcgttagcttttctagcccccttcggtggtgttc
1 M S G P V P N F P Y K L F N I P L A L A F L A P S V V F
3311 tttacttcgatccatttatcgatccagcctttgaacatatcacaagaagctttgaacatatccgtaa 3243
29 F T S I H L S I Q P L N I S Q E A L N I Y P *
182ORF049
1578 atgttgcaatctcaagagcgcaaatcaaagaagcgcaaatataaacagagcaagctcaaaaagcgaaagaagaactacaaag
1 M L Q S Q E R K S K K R K L K Q S K L K K R K K N T T K
1662 agcttaacaagttgaagttgaagccacagaaaacacaattgtcacccaactattttaa 1724
29 S L T K L K L R S P Q K T Q L S H Q L F *
182ORF050
8012 atggttatcttgggtttctttaaagacctacacttgggttcattggtttgcgaggggcagaagatggtcaaactcgatcattatc
1 M V I L V S L K T L H L G S W F A Q G Q K M V K S I I I
8096 acaaccttattttctttacagcaaacgaagcaatgtatcacaagagatatcctgttttaa 8155
29 T T L F S L Q Q T K Q C I T R D I L F *
182ORF051
9390 atgcttctgaaaaagaacaagaacacagacattaataaagatcaaaatcaaaccaagatacgattacacgatataaaggta
1 M L L K K K Q R T Q T L I K I K I K P K I R L H D I K V
9474 aaaaggaacactgattatgctgacttactcgaaaatatcgtagaagtggtttga 9530
29 K R E T L I M L T Y S K N I V E V F *
182ORF052
4096 gtgatagttgacaagagtcataatttggcagattggcgcaatgtacacgtgaaatatcggtcgctcccgttaagttatggacac
1 V I V D K S Q I W R D W A N V H V K Y R A L P L S Y G H
4180 ataaacgttttgaccgtcaaccaatcgcaaaaaccttttaggagtagcccttaa 4233
29 I N V L T V N Q S Q K P F R S S P *
182ORF053
15656 gtggaacagaatacgaagatttttagtatcaacaatgagtttgcataatgatagtgatacgttttattgaaaagagaagtaaaata
1 V E Q N T K I L V S T M S L S M I V I R L L K R E V K I
15740 gtagtttcttatgcccattgcttttgaagggaatactttgggtattggatag 15793
29 V V S Y A P L L L K G K S L G I G *
182ORF054
8136 gtgatacattgcttctgttgaagaaatagggttgataatgatcgatttgaccatcttctgcccctgcgcaaacat
1 V I H C F V C C K E N R V V I M I D L T I F C P C A N H
8052 gaacccaagtgtagggtctttaagaacaaagataaccattagtggttaa 8002
29 E P K C R V F K E T K I T I S V *
182ORF055
8324 atgaaaaaataacttctcattgctacaagcttataaccaaatgacgaaaataatcaggctgtttttgtggataaagatatgg
1 M K R N T S H C Y K L I T K L T K I I R L F L W I K I W
8408 agtttgacgaatcttttaattgtatggcaacaaatgctccatatgtag 8455
29 S L T N L L M Y G K Q M L H M *
182ORF056
6549 gtggcccatctccttttctattatttacttctcatcaattcaagtggggagggtatacaaaccaaatggggcaggcaatgcta
1 V A H L L F P I I Y F L S I Q V G R Y T N Q M G Q A M L
6633 attttggagagtacatggcgtttcttacaacgaagaaccttttttaa 6680
29 I L E S T W R F L Q R K N L F *
182ORF057
8264 atgtccgccatctctaaagcaaacgatgtaaaacttggttaacgtaggaactttcaagtcattattatacaacatgatactttt
1 M S A I S K A K R C K L G N V G T F K S L L Y N M I H F
8180 gatttatcatcatcatcatcatatcttaaacaggatattctcttgta 8133
29 D L S S S S S Y L K T G Y L L *
182ORF058
5176 gtgtattcaaatctgcttacttctgacacctgtgtataaagcgttcattacaccagcaacgaactattgaaattatcccatgaa
1 V Y S N S L T S S P V Y K A F I T P A T K L L K L S H E
5092 gtaaatgctttttctaacatgcttcttggtcggtttgtttgtag 5048
29 V N A F S N H A S W I V C L *
182ORF059
15876 atggtctttctgtagtcattgcatataaaatgatttggatttggatgataactcacatagacacaacctgtttcagcgtc
1 M V F R S H C I K M I C I W L I I I T H I D T T C F S V
15792 tatccaatacccaagattttcccttcaaaagcaatggcgcataa 15748
29 Y P I P K D F P F K S N G A *
182ORF060
15404 gtgatttttgattttctcaattaaaaactcatcaacaaaattgtacgaacttcgggatattcattagatttttcaattccccac
1 V I F D F S I K N S S N K I V R T S G Y S L D F S I P H
15320 gtactaagtggacagcccaacccattatttatcatcacaatag 15276
29 V L S G T A Q P I N L S S Q *
182ORF061
2102 atgaggggacttctccacctgtttcagactcgatcacttttgcaatcttactgtaaacttgttctttttctgtgtacttctg

327

1 M R G L L H L F Q T R S L L Q S Y C K L V L F S V V L L
2018 cttcgtcataaatgtagtcaagggttcattgcttaagaagtactaa 1974
29 L R H K C S Q G S C L R S Y *
182ORF062
1992 atgtctaagaagtactaacaatggttttcataaataagatcaagccccattgagtcagtaaacgaacaatatcgtagcgtct
1 M S K K L L T Y V F I N R S S P I E S S K R T I S S A S
1908 gaattgaaaatagtaaacatcgcttctgctgaattgtcgtaa 1867
29 E L K I V N I A C L K L S *
182ORF063
14306 gtgtaccttctaaccctctcatgcgcaaaatgatacacaccaatctttttacctaagacaaaagcttggtaaagtgcggt
1 V Y L L N P S H A Q N D T H Q S F Y L K T K L V E M L G
14222 cacaatcaggggtttacataaacctgttcgcgctgttgccttaa 14181
29 H N Q G L H N L F R L L L *
182ORF064
7356 atgatgttagtcaaaccaaaaagggttgttactgtgaaggctgaaaagatcgctcctctgtactcattgcactgtttccc
1 M M L V K P T K G L L L A K A E K I A P P V L I A L F P
7272 ataccatgtctgaaagtattgcaagtgtttgtctctga 7234
29 I P C L K V L R M F C S *
182ORF065
3582 atgaatgctatctgtatcacataaataatgcgatcaaaacatttttgagcgggtgtaaggtagtatatctacccaagccgt
1 M N A I C I T I N N A I K T F L S G C N G S I S T P S R
3498 caaaaactagcaagcggaacataaacaggatctcttaa 3460
29 H K T S K R N I N R I S *
182ORF066
4234 atgtggctactctttttgtgtttcacagaattatgtttcacgtgaacagtttttatggataataagaatcaaaaggagggtg
1 M W L L F F V F H R I M F H V K Q F L W Y N R I K R R W
4318 agattatggaataaagaacatgaatcaatttttaa 4353
29 R L W K L K N M N Q F *
182ORF067
13882 atgatactgcttagctttaaaactactaaagtcgatttctctgttaaatgttgcaattgtaaaacctaacacaaaatcgata
1 M I P A L A L K L L K S I S L L N L A I V K P N T K S I
13798 atcattgcaaccattaaccatataatcaaacataa 13763
29 I I A T I N H I I K P *
182ORF068
7267 atgtctgaaagtattgcaagtgtttgtctttagcaatcaaggagttttgtttccttgcatgaatgcagaagcatagtcaga
1 M S E S I A N V L L L S N Q G V F V S L H E C R S I V R
7183 tttaactcctacatcgtaggatcattatcgattaa 7148
29 F N S Y I V R I I I D *
182ORF069
5027 gtggaacaatgtttttacatcggaacttctgttttaataacctctgttaacagactcgtagcagggtgaaacttatgttctctgtgc
1 V E Q C F Y I G N F L F K Y P C N R L V R V E L M F L C
4943 aatgtcaacaaaaatttcttcaatcggttcgacctaa 4908
29 N V N K N F F N R S T *
182ORF070
1031 gtgatgggttcggtccacaaaaccagaaactcgtctgagtaaaactagaatatctttcaattctagtagtctcgccaattgct
1 V M V R L H Q N Q K L R L S K L E Y L S I L V L R Q L R
947 tacgtaacggaattgaagccccgtttgtggcattga 912
29 Y V T E L K P R L W H *
182ORF071
11741 atgggtttgcattatggttgccacaaggcgctcaaagtggttaaagggaattttctttaatgatactcgcaattacaatcggtttg
1 M V L H Y G C H K A L K V V K E F S L M I L A I T I V L
11825 actttgattgtttgttcgtaactgtactttaa 11857
29 T L I C L F V T V L *
182ORF072
11723 atgtttacattaaatgcgcgtcattgtttcaactttaatgtcgtttctcccgatcctaagaaagtaactacaggtacatcacgt
1 M F T L N A V I V S N F N V V S P D P K K V T T G T S R
11639 ttcaattcaatgtgttagcaagcgataa 11610
29 F N S M V L A K R *
182ORF073
2876 gtgaagccgctttgtatgctttacgtaagtctttatcaaaccttaaagacaaaataggaaccattgtttgaaagttgatttt
1 V K P P L Y A L R K S L S N P K D K I G N H C L K V D F
2792 ccatgtgtagcttttagccaatctttgtaa 2763
29 P C V A P S Q S L *
182ORF074
8923 gtgattgataaattttgtttcaaatctgctcgtttgtttcgtcataaaacggataatcaaaatcaacaattgttttcgccc
1 V I D K F C F K F C S F C F V I K R I I K I K Q L F S A
8839 aacttcaatcgtttcttttcgagataa 8813
29 N F N T F F S R *
182ORF075
7463 gtgttacattatctggaatattttcgatatctgccactttacgtgccaagaggttcaaaccgttttttttcagaaacatagt
1 V L H Y L E Y F R Y L P L Y L P R G S N R F L F Q K H S
7379 tgtttactgtgtcctgctcccatga 7353
29 C L L V V L L P *
182ORF076
2426 atgagtggtggaagtgttcgatcttctttgttctttacaccatttgaaaccatttttgataaccatgaaagcataaactct

328

1 M S V E N V R S S F A S L H H L K P F L N N H E S I N S
2342 ccgtcaaatttttcgttgaggaaataa 2316
29 P S N F S L W K *
182ORF077
11858 atgaaggaacgtatgttggttgctagaggtagaggggttacatttgaaaattgtctattctctaataatctctcaagcaatta
1 M K E R M L L L L E V E G L H L K I V Y S L I S L K Q L
11942 tcaaaacagcttttcccgatgtaa 11965
29 S K Q L F P M *
182ORF078
7671 gtgcctacaatatatttggttcttttaatttaataatgaaattccatgcttttcttggttgtaagtttggtgtagctactcgattgctc
1 V P T I F G S F N L M K F H A F L V C K F G V A T R L L
7587 tttgtgccatacattgagaagtaa 7564
29 F V P Y I E K *
182ORF079
7488 gtgaaagataagtttgatccaagctgtgttacattatctggaatattttcgatatctgccactttacctgccagagggttcaaa
1 V K D K F D P S C V T L S G I F S I S A T L P A K R F K
7404 ccgttttcttttccagaaacatag 7381
29 P F S F S E T *
182ORF080
4473 gtgtgctatttgctgatgtcaaagcttcagttgttgctccgtagtcttctcgcaatgcttcaagatgttctacaatctttgatc
1 V C Y L L M S K L Q L L L R S L L A M L Q D V L Q S L I
4389 ttgcttcaccgtctgtga 4372
29 L L H R L *

Table 24

Sequence similarities phage 182 and public databases

Phage: 182

Database: nr

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2
(604 letters)

gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9) >...	384	e-105
gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9) >...	374	e-103
gi 1429238 gnl PID e1173412 (X99260) tail protein [Bacteriophag...	346	3e-94
gi 215339 (M12456) p9 tail protein [Bacteriophage phi-29] >gi 2...	208	8e-53
gi 1181970 gnl PID e221269 (Z47794) tail protein [Bacteriophage...	62	8e-09
gi 1181968 gnl PID e221267 (Z47794) tail protein [Bacteriophage...	56	6e-07
gi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM...	49	8e-05

Query= sid|110157|lan|182ORF002 Phage 182 ORF|2152-3873|1
(573 letters)

gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE >gi 76896 pir JQ0...	665	0.0
gi 1429230 gnl PID e1173404 (X99260) DNA polymerase [Bacterioph...	657	0.0
gi 118849 sp P03680 DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP...	654	0.0
gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP...	654	0.0
gi 15732 (X53371) DNA polymerase (AA 1-575) [Bacteriophage phi-29]	651	0.0
gi 15734 (X53370) DNA polymerase (AA 1-575) [Bacteriophage phi-29]	651	0.0
gi 1572479 gnl PID e242301 (X96987) DNA polymerase [Bacterioph...	565	e-160
gi 1072656 pir S51275 DNA polymerase - phage CP-1 >gi 836593 g...	301	1e-80
gi 118847 sp P22374 DPOM_ASCIM PROBABLE DNA POLYMERASE >gi 8385...	71	3e-11
gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE >gi 2833...	65	1e-09
gi 461963 sp P33538 DPOM_NEUIN PROBABLE DNA POLYMERASE >gi 1018...	62	1e-08
gi 1084487 pir S41618 DNA polymerase - slime mold (Physarum po...	61	3e-08
gi 2435429 (AF012250) unassigned reading frame (possible DNA po...	61	3e-08
gi 578157 gnl PID e246743 (X52106) DNA polymerase [Neurospora i...	59	1e-07
gi 2147969 pir S72369 probable DNA-polymerase - Gelasinospora ...	58	2e-07
gi 2147968 pir S62752 probable DNA-polymerase - Gelasinospora ...	58	2e-07
gi 3511140 (AF061244) B type DNA polymerase [Agrocyebe aegerita]	57	3e-07
gi 118850 sp P10479 DPOL_BPPRD DNA POLYMERASE (PROTEIN P1) >gi ...	56	6e-07
gi 578144 (X63909) putative DNA-polymerase, B-type [Morchella c...	47	3e-04
gi 232013 sp P30322 DPOM_AGABT PROBABLE DNA POLYMERASE >gi 3208...	46	6e-04

Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
(442 letters)

gi 138117 sp P13849 VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ...	309	2e-83
gi 138118 sp P07531 VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ...	305	3e-82
gi 1429236 gnl PID e1173410 (X99260) major head protein [Bacter...	300	1e-80
gi 1181958 gnl PID e221257 (Z47794) major head protein [Bacteri...	152	6e-36

Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
(349 letters)

gi 137932 sp P15132 VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR...	52	8e-06
gi 1429242 gnl PID e1173416 (X99260) morphogenesis protein [Bac...	48	7e-05
gi 137933 sp P07538 VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR...	47	2e-04

Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
(343 letters)

gi 137944 sp P11014 VG16_BPPH2 ENCAPSIDATION PROTEIN (LATE PROT...	402	e-111
gi 137945 sp P07541 VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROT...	402	e-111
gi 1429245 gnl PID e1173419 (X99260) encapsidation protein [Bac...	381	e-105
gi 1181972 gnl PID e221271 (Z47794) encapsidation protein [Bact...	159	2e-38

Query= sid|110162|lan|182ORF007 Phage 182 ORF|7795-8775|1
(326 letters)

gi 1429239 gnl PID e1173413 (X99260) upper collar protein [Bact...	271	5e-72
gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...	256	1e-67
gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...	256	2e-67
gi 1181960 gnl PID e221259 (Z47794) connector protein [Bacterio...	148	6e-35

Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
(292 letters)

gi 4210750 gnl PID e1374037 (AJ132604) LysL protein [Lactococcu...	139	2e-32
gi 462559 sp P34020 LYC_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC...	75	8e-13
gi 2327014 (U82823) putative lysozyme [Saccharopolyspora erythr...	64	2e-09
gi 126652 sp P25310 LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA-...	60	2e-08
gi 127789 sp P19386 LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	60	2e-08
gi 67761 pir MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5...	59	3e-08
gi 4105636 (AF049087) lys [Leuconostoc oenos bacteriophage 10MC]	59	3e-08
gi 623084 (L02496) muramidase; muramidase [Bacteriophage LL-H]	57	1e-07
gi 127787 sp P15057 LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	2e-07
gi 126597 sp P00721 LYCH_CHASP N,O-DIACETYLMURAMIDASE (LYSOZYME...	57	2e-07
gi 127788 sp P19385 LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	2e-07
gi 67762 pir MUBPC7 N-acetylmuramoyl-L-alanine amidase (EC 3.5...	56	3e-07
gi 3025168 sp P76421 YEGX_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN...	53	2e-06
gi 4204413 (AF047001) Lys44 [Oenococcus oeni temperate bacterio...	53	3e-06
gi 2116978 gnl PID d1020940 (D88151) cortical fragment-lytic en...	52	5e-06
gi 2392844 (AF011378) lysin [Bacteriophage sk1]	48	8e-05

Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
(278 letters)

gi 1429240 gnl PID e1173414 (X99260) lower collar protein [Bact...	180	1e-44
gi 137921 sp P04333 VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE...	171	5e-42
gi 215341 (M12456) p11 lower collar protein [Bacteriophage phi-29]	98	9e-20
gi 224162 prf 1011232B protein p11, lower collar [Bacteriophage...	97	1e-19
gi 535260 (Z30339) STARP antigen [Plasmodium reichenowi]	50	1e-05
gi 4049753 (AF063866) ORF MSV230 hypothetical protein [Melanopl...	49	4e-05
gi 2131557 pir S70306 hypothetical protein YEL077c - yeast (Sa...	48	5e-05
gi 131782 sp P12753 RAS0_YEAST DNA REPAIR PROTEIN RAD50 (153 KD...	48	7e-05
gi 2131309 pir S70305 hypothetical protein YBL113c - yeast (Sa...	47	2e-04
gi 499325 (Z26314) STARP antigen [Plasmodium falciparum]	46	3e-04
gi 3845171 (AE001391) ribosome releasing factor (OO, TP) [Plasm...	46	3e-04
gi 731903 sp P40434 YIR7_YEAST HYPOTHETICAL 197.5 KD PROTEIN IN...	45	5e-04
gi 1632829 gnl PID e276379 (Y08924) AARP2 protein [Plasmodium f...	45	5e-04
gi 1176490 sp P40889 YJW5_YEAST HYPOTHETICAL 197.6 KD PROTEIN I...	45	5e-04
gi 1077300 pir S51848 hypothetical protein HRD1054 - yeast (Sa...	45	5e-04
gi 2425143 (AF020407) Wima [Dictyostelium discoideum]	45	6e-04
gi 1181961 gnl PID e221260 (Z47794) collar protein [Bacterioph...	45	6e-04
gi 2132657 pir S64819 probable membrane protein YLL067c - yeas...	45	8e-04
gi 2133041 pir S65341 probable membrane protein YPR204w - yeas...	45	8e-04
gi 730275 sp P39793 PBPA_BACSU PENICILLIN-BINDING PROTEINS 1A/1...	45	8e-04

Query= sid|110165|lan|182ORF010 Phage 182 ORF|1310-2155|2
(281 letters)

gi 135604 sp P06812 TERM_BPNF DNA TERMINAL PROTEIN >gi 75815 pi...	69	3e-11
gi 1572478 gnl PID e242334 (X96987) terminal protein [Bacteriop...	65	3e-10
gi 1429231 gnl PID e1173405 (X99260) terminal protein [Bacterio...	64	1e-09

Query= sid|110166|lan|182ORF011 Phage 182 ORF|9607-10158|1
(183 letters)

gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE...	51	6e-06
gi 1429241 gnl PID e1173415 (X99260) pre-neck appendage protein...	51	6e-06
gi 137927 sp P20345 VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE...	50	1e-05

Query= sid|110169|lan|182ORF014 Phage 182 ORF|13716-14108|3
(130 letters)

gi 137936 sp P11188 VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14...	97	6e-20
gi 137938 sp P07539 VG14_BPPZA LYSIS PROTEIN (LATE PROTEIN GP14...	96	8e-20
gi 1429243 gnl PID e1173417 (X99260) lysis protein [Bacterioph...	96	8e-20
gi 215332 (M14782) lysis protein [Bacteriophage phi-29]	94	5e-19

Query= sid|110170|lan|182ORF015 Phage 182 ORF|854-1225|2
(123 letters)

331

gi|15670 (V01155) reading frame 10 (may be gene 4) [Bacterioph... 70 5e-12
 gi|138072|sp|P06953|VG5A_BPPZA EARLY PROTEIN GP5A >gi|75836|pir... 69 7e-12

Query= sid|110174|lan|182ORF019 Phage 182 ORF|4323-4613|3
 (96 letters)

gi|1429235|gnl|PID|e1173409 (X99260) head morphogenesis protein... 61 2e-09
 gi|138111|sp|P13848|VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE ... 57 3e-08
 gi|138112|sp|P07533|VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE ... 54 1e-07

Query= sid|110180|lan|182ORF025 Phage 182 ORF|548-814|2
 (88 letters)

gi|138099|sp|P06955|VG6_BPPZA EARLY PROTEIN GP6 >gi|75841|pir|... 55 7e-08
 gi|138098|sp|P03685|VG6_BPPH2 EARLY PROTEIN GP6 >gi|75840|pir|... 54 2e-07
 gi|1429234|gnl|PID|e1173408 (X99260) gene 6 product [Bacterioph... 54 2e-07

Table 25

Homologies between 182 ORFs and proteins in public databases

Phage: 182

Database: Swissprot

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2
(604 letters)

gi 138124 sp P07534 VG9_BPPZA TAIL PROTEIN (LATE PROTEIN GP9)	384	e-106
gi 138123 sp P04331 VG9_BPPH2 TAIL PROTEIN (LATE PROTEIN GP9)	374	e-103
gi 2500030 sp Q59968 CARA_SULSO CARBAMOYL-PHOSPHATE SYNTHASE SM...	49	2e-05

Query= sid|110157|lan|182ORF002 Phage 182 ORF|2152-3873|1
(573 letters)

gi 118848 sp P19894 DPOL_BPM2 DNA POLYMERASE	665	0.0
gi 118849 sp P03680 DPOL_BPPH2 DNA POLYMERASE (EARLY PROTEIN GP2)	654	0.0
gi 118851 sp P06950 DPOL_BPPZA DNA POLYMERASE (EARLY PROTEIN GP2)	654	0.0
gi 118847 sp P22374 DPOM_ASCIM PROBABLE DNA POLYMERASE	71	7e-12
gi 461962 sp P33537 DPOM_NEUCR PROBABLE DNA POLYMERASE	65	3e-10
gi 461963 sp P33538 DPOM_NEUIN PROBABLE DNA POLYMERASE	62	3e-09
gi 118850 sp P10479 DPOL_BPPRD DNA POLYMERASE (PROTEIN P1)	56	2e-07
gi 232013 sp P30322 DPOM_AGABT PROBABLE DNA POLYMERASE	46	2e-04
gi 118887 sp P10582 DPOM_MAIZE DNA POLYMERASE (S-1 DNA ORF 3)	46	2e-04

Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
(442 letters)

gi 138117 sp P13849 VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN ...	309	6e-84
gi 138118 sp P07531 VG8_BPPZA MAJOR HEAD PROTEIN (LATE PROTEIN ...	305	7e-83

Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
(349 letters)

gi 137932 sp P15132 VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE PR...	52	2e-06
gi 137933 sp P07538 VG13_BPPZA MORPHOGENESIS PROTEIN 1 (LATE PR...	47	6e-05

Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
(343 letters)

gi 137945 sp P07541 VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROT...	402	e-112
gi 137944 sp P11014 VG16_BPPH2 ENCAPSIDATION PROTEIN (LATE PROT...	402	e-112

Query= sid|110162|lan|182ORF007 Phage 182 ORF|7795-8775|1
(326 letters)

gi 137915 sp P07535 VG10_BPPZA UPPER COLLAR PROTEIN (CONNECTOR ...	256	3e-68
gi 137914 sp P04332 VG10_BPPH2 UPPER COLLAR PROTEIN (CONNECTOR ...	256	5e-68

Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
(292 letters)

gi 462559 sp P34020 LYC_CLOAB AUTOLYTIC LYSOZYME (1,4-BETA-N-AC...	75	2e-13
gi 126652 sp P25310 LYCM_STRGL LYSOZYME M1 PRECURSOR (1,4-BETA-...	60	5e-09
gi 127789 sp P19386 LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	60	5e-09
gi 127787 sp P15057 LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	4e-08
gi 126597 sp P00721 LYCH_CHASP N,O-DIACETILMURAMIDASE (LYSOZYME...	57	4e-08
gi 127788 sp P19385 LYCA_BPCP7 LYSOZYME (ENDOLYSIN) (MURAMIDASE...	57	5e-08
gi 3025168 sp P76421 YEGX_ECOLI HYPOTHETICAL 32.0 KD PROTEIN IN...	53	5e-07

Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
(278 letters)

gi 137921 sp P04333 VG11_BPPH2 LOWER COLLAR PROTEIN (LATE PROTE...	171	1e-42
gi 131782 sp P12753 RA50_YEAST DNA REPAIR PROTEIN RAD50 (153 KD...	48	2e-05
gi 1176490 sp P40889 YJW5_YEAST HYPOTHETICAL 197.6 KD PROTEIN I...	45	1e-04
gi 731903 sp P40434 YIR7_YEAST HYPOTHETICAL 197.5 KD PROTEIN IN...	45	1e-04
gi 730275 sp P39793 PBPA_BACSU PENICILLIN-BINDING PROTEINS 1A/1...	45	2e-04
gi 1168610 sp P41696 AZF1_YEAST ASPARAGINE-RICH ZINC FINGER PRO...	44	3e-04

333

gi 731587 sp P38900 YH19_YEAST HYPOTHETICAL 70.1 KD PROTEIN IN ...	44	3e-04
Query= sid 110165 lan 182ORF010 Phage 182 ORF 1310-2155 2 (281 letters)		
gi 135604 sp P06812 TERM_BPNF DNA TERMINAL PROTEIN	69	8e-12
Query= sid 110166 lan 182ORF011 Phage 182 ORF 9607-10158 1 (183 letters)		
gi 137928 sp P07537 VG12_BPPZA PRE-NECK APPENDAGE PROTEIN (LATE...	51	2e-06
gi 137927 sp P20345 VG12_BPPH2 PRE-NECK APPENDAGE PROTEIN (LATE...	50	3e-06
Query= sid 110169 lan 182ORF014 Phage 182 ORF 13716-14108 3 (130 letters)		
gi 137936 sp P11188 VG14_BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)	97	2e-20
gi 137938 sp P07539 VG14_BPPZA LYSIS PROTEIN (LATE PROTEIN GP14)	96	2e-20
Query= sid 110170 lan 182ORF015 Phage 182 ORF 854-1225 2 (123 letters)		
gi 138072 sp P06953 VG5A_BPPZA EARLY PROTEIN GP5A	69	2e-12
Query= sid 110174 lan 182ORF019 Phage 182 ORF 4323-4613 3 (96 letters)		
gi 138111 sp P13848 VG7_BPPH2 HEAD MORPHOGENESIS PROTEIN (LATE ...	57	9e-09
gi 138112 sp P07533 VG7_BPPZA HEAD MORPHOGENESIS PROTEIN (LATE ...	54	4e-08
Query= sid 110180 lan 182ORF025 Phage 182 ORF 548-814 2 (88 letters)		
gi 138099 sp P06955 VG6_BPPZA EARLY PROTEIN GP6	55	2e-08
gi 138098 sp P03685 VG6_BPPH2 EARLY PROTEIN GP6	54	5e-08

334

BLASTP 2.0.8 [Jan-05-1999]

Query= sid|110156|lan|182ORF001 Phage 182 ORF|5966-7780|2
(604 letters)

>gi|138124|sp|P07534|VG9_BPPZA_TAIL_PROTEIN (LATE PROTEIN GP9)
>gi|75849|pir|WMBP9Z gene 9 protein - phage PZA
>gi|216058 (M11813) tail protein [Bacteriophage PZA]
Length = 599

Score = 384 bits (975), Expect = e-105

Identities = 231/610 (37%), Positives = 344/610 (55%), Gaps = 36/610 (5%)

Query: 6 TNVLLANVPFDNTYTHTRWFKTQQEQESYFNSFPVLNENRDCSYQRDTQLGGVFRVDKH 65
TNV++LA+VPP N Y +TRWF + Q ++FNS + E ++Q + V
Sbjct: 9 TNVRILADVPPSNDYKNTRWFTSSSNQYNWFNSKTRVYEMSKVTFQGFRENKSYISVSLR 68

Query: 66 KDALYACNYLIFKNEETYPKQYAFVTDIEYKNDNTSFVTFEIDVLQTYRFDIGIRESF 125
D LY +Y++F+N + Y +KW YAFVT++EYKN T++V FEIDVLQT+ F+I +ESF
Sbjct: 69 LDLLYNASYIMFQAD-YGNKWFYAFVTELEYKQVGT+TVVHFEIDVLQTMFMNKFQESF 127

Query: 126 IAKEHPQLYYSNGIPIINTIESLDYGREYTTTNVITTFHPNDGVNFLVILTSEAM--PVG 183
I +EH +L+ +G P INTI+E L+YG EY +V P D + FLV+++ M G
Sbjct: 128 IVREHVKLWDDGTPTINTIDEGLNYGSEYDIVSVENHRYDDMMFLVVISKSIMHGTAG 187

Query: 184 DKEDKSG---GSIVGGSPSPFSYLLPINSSGEVYKPN-GAGNANFGEYMAFLT---TKEP 236
+ E + S+ G P P YY+ P G+V K G NAN + LT +++
Sbjct: 188 EAESRLNDINASLNGMPQLCYIHPFYKDGKVPKTFIGDNNANLSPIVNMLTNIFSQKS 247

Query: 237 FLNKIVGMVYTSYTGIPFIVDHANKTVRYNAGGSYKIMLPTYASDPTGMTKFAFFCVKE 296
+N IV MYVT Y G+ + +K ++ + + A D G + T VK+
Sbjct: 248 AVNNIVNMYVTDYIGLKLDDYKNGDKELKLDKDMFEQAGI---ADKKGNVDTIF---VKK 301

Query: 297 ARTFVFKRIDLVGNVYNYFREAFPFNVKESKLFMYPYCLIEITDTKGHVMTLRPEYLTGG 356
+ ID G+ + F + +ESKL MYPYC+ E+TD KG+ M L+ EY+
Sbjct: 302 IPDYETLEID-TGDKWGGFTKD-----QESKLMMYPYCVTEVTDKFGNHMNLKTEYIDNN 355

Query: 357 KLSVYVKGSLGISNKMVIEPIDYDVSNTI----ITNLSKMLIDNDPNDVGKSDYASA 412
KL + V+GSLG+SNKV DY+ S +T D LI+N+PND+ + +DY SA
Sbjct: 356 KLKIQVRGSLGVSNNKVAYSIQDYNAGGSLGGDRLTASLDTSLINNPNNDIAIINDYLSA 415

Query: 413 FMQGNKNSLIAEQNIRNTFRHGMNSAMSTGGAIFSAASNNPFVGLTNIMGAGQQVNN 472
++QGNKNSL Q+ +I GM +S G ++ +PF +++ G N
Sbjct: 416 YLQGNKNSLENQSSILFNGIVGMLGGVSAG----ASAVGRSPFGLASSVTGMTSTAGN 471

Query: 473 YVSEKENGLNLLAGKVADIENIPDNVTQLGSNLSFTTGN-FQNYQLRFPKQIKYEYATRL 531
V + + L K ADI NIP +T++G N +F GN ++ Y ++ KQ+K EY L
Sbjct: 472 AVLDD---MQALQAKQADIANIPPQLTKMGNTAFDYGNGYRGVYVIK-KQLKAEYRRSL 526

Query: 532 DRYFSMYGTKSNRVATPNLQTRKAWNFIKLKEPNVGTMSNDVLTRVKQIFSAGVTLWHT 591
+F YG K NRV PNL+TRKA+N+I+ K+ I G ++N+ L ++ IF G+TLWHT
Sbjct: 527 SSFFHKYGYKINRVKKNLRTKAYNYIQTKDCFISGDINNNDLQEI RTIFDNGITLWHT 586

Query: 592 NDVLNYNQDN 601
+D+ NY+ +N
Sbjct: 587 DDIGNYSVEN 596

Query= sid|110157|lan|182ORF002 Phage 182 ORF|2152-3873|1
(573 letters)

>gi|118848|sp|P19894|DPOL_BPM2_DNA_POLYMERASE >gi|76896|pir|JQ0161
DNA-directed DNA polymerase (EC 2.7.7.7) - phage M2
>gi|215509 (M33144) DNA polymerase [Bacteriophage M2]
Length = 572

Score = 665 bits (1697), Expect = 0.0

Identities = 327/589 (55%), Positives = 420/589 (70%), Gaps = 38/589 (6%)

Query: 3 KKYTGDFETTTDLNDCRVWSWGVCDIDNVNMTFGLIDSFFEWCKMQGSTDIYFHNK 62
K ++ DFETTT L+DCRVW++G +I N+DN G +D F +W M+ D+YFHN KF

335

Sbjct: 4 KMFSCDFETTTKLDDCRVWAYGYMEIGNLDNYKIGNSLDEFMQWV-MEIQADLYFHNLF 62

Query: 63 DGEFMLSFLKNGFKWCKEAKEDRTFSTLISNMGQWYALEICWEVNYXXXXXXXXXXXXX 122
 DG F+++WL ++GFKW E + T++T+IS MGQWY ++IC+

Sbjct: 63 DGAFIVNWLEQHGFKWSNEGLPN-TYNTIISKMGQWYMIDICFGYK-----GKRKL 112

Query: 123 XXIIYDSLKKYPFPVKQIAEAFNFPKKGIDYTKERPIGYKPTKDEWEYLKNDIQIMAM 182
 +IYDSLKK PFPVK+IA+ F P+ KG+IDY ERP+G++ T +E+EY+KNDI+I+A

Sbjct: 113 HTVIYDSLKKLPFPVKIAKDFQLPLKGDIDYHTERPVGHEITPEEYKNDIEIAR 172

Query: 183 ALKIQFDQGLTRMTRGSDALGDYKDWLKAHKGSTFKQWFPILSLGFDKDLRKAYKGGFT 242
 AL IQF QGL RMT GSD+L +KD L F + FP LSL DK++RKAY+GGFT

Sbjct: 173 ALDIQFKQGLDRMTAGSDSLKGFKDILST----KKFNKVPKLSLPMDEIRKAYRGGFT 228

Query: 243 WVNKVFQKGEIGDGVFDVNSLYPSQMYRPLPYGTPLFYEGEYKPNNDYPLYIQNIKVR 302
 W+N ++ KEIG+G+VFDVNSLYPSQMY RPLPYG P+ ++G+Y+ + YPLYIQ I+

Sbjct: 229 WLNDKYKEKEIGEMVFDVNSLYPSQMYRPLPYGAPIVFGKYEKDEQYPLYIQRIRFE 288

Query: 303 FRLKEGYIPTIQVKQSSLFIQNEYLESSVNLGVDELIDLTNTVDLELFFEHYDILEIH 362
 F LKEGYIPTIQ+K++ F NEYL++S GV E ++L LTNVDELE EHY++ +

Sbjct: 289 FELKEGYIPTIQIKKNPFFKGYEYLNKNS----GV-EPVELYLTNVDELELIQEYELYNVE 343

Query: 363 YTYGYMFKASCDMFKGWIDKWEVKNTTEGARKANAKGMLNSLYGKFGTNPDTGKVPYM 422
 Y G+ F+ +FK +IDKW VK EGA+K AK MLNSLYGKF +NPD+TGKVPY+

Sbjct: 344 YIDGFKFRBKTLGKDFIDKNTYVKTHEEGAKQLAKLMLNSLYGKFASNPDTGKVPYL 403

Query: 423 GEDGIVRLTLGEEELRDPVYVPLASFVTAWGRYTTITTAQKCFDRIIYCDTDSIHLVGT 482
 +DG + +G+EE +DPVY P+ F+TAW R+TTIT AQ C+DRIIYCDTDSIHL GTE

Sbjct: 404 KDDGSLGFRVGDDEYKDPVYTPMGVFITAWARFTTITAAQCYDRIIYCDTDSIHLTGT 463

Query: 483 VPEAIDHLVDPKLLGYWGHESTFQRAKFIQKT-----YVEEIDGEL----- 524
 VPE I +VDPKLLGYW HESTF+RAK++RQKT YV+E+DG+L

Sbjct: 464 VPEIKDIVDPKLLGYWAHESTFKRAKYLQKTYIQDIYVKEVDGKLKECSPDEATTTKF 523

Query: 525 NVKCAGMPDRIKEIVTFDNFEVGFSSYGKLLPKRTQGGVVLVDTMFTIK 573
 +VKACGM D IK+ VTFDNF VGFSS GK P + GGVVLVD++FTIK

Sbjct: 524 SVKCAGMTDTIKKVTFDNFVAVGFSSMGKPKPVQVNGGVVLVDSVFTIK 572

Query= sid|110159|lan|182ORF004 Phage 182 ORF|4626-5954|3
 (442 letters)

>gi|138117|sp|P13849|VG8_BPPH2 MAJOR HEAD PROTEIN (LATE PROTEIN GP8)
 >gi|75845|pir||WMBP89 gene 8 protein - phage phi-29
 >gi|215325 (M14782) major head protein [Bacteriophage
 phi-29] >gi|225362|prf||13012708 gene 8 [Bacillus sp.]
 Length = 448

Score = 309 bits (783), Expect = 2e-83
 Identities = 176/440 (40%), Positives = 250/440 (56%), Gaps = 27/440 (6%)

Query: 4 KITEQDVLRLATNVETPVQLMTAIYNSSSSLFQANVMPNADNIEAVGAGITRLDVVKNEF 63
 ++ AI NS F++ VP+ A+N+ VGAGI V+N+F

Sbjct: 2 RITFNDVKTSLGITESYDIVNAIRNSQGDNFKSYVPLATANNVAEVGAGILINQTVQNDP 61

Query: 64 ISTLVDRIGKVIRYKSWRNPLKMFKKGNMPLGRTIEEIFVDIAQEHKFPNDESVTGVFK 123
 I++LVDRIG VVIR S NPLK FKKG +PLGRTIEEI+ DI +E +++ +E+ VF+

Sbjct: 62 ITSLVDRIGLVVIRQVSLNPLKFKKQIPLGRTIEEITYDITKEKQYDAEAEQKVFE 121

Query: 124 QEVPDVKTLFHEINREGYYKTIQEAWLEKAFSTWDFNSFVAGVMNALYTGDEVSEFEY 183
 +E+P+VKTLFHE NR+G+Y QTIQ+ L+ AF SW NF SPV+ ++NA+Y EV E+EY

Sbjct: 122 REMPNVKTLFHERNRQGFYHQTIQDDSLKTAFAVSWGNFESFVSSIINAIYNSAEVDEY 181

Query: 184 TKLLIANYQEKELFKEIEIGETESNA--KEFIRKIKSTSNKLEFM--SSAYNAQGVKTS 239
 KLL+ NY K LF ++I E T S EF++K+++T+ KL S +N+ V+T

Sbjct: 182 MKLLVDNYYSKGLFTTVKIDEPTSSGTALTEFVKMRATARKLTLPGQSRDWNMAVRTR 241

Query: 240 TSKSDQYXXXXXXXXXXXXXXXXXXXXXFNMSKTDVFGHKIVIDEFPKKEGEESNIVAVIV 299
 + D + FNM++TDF+G+ VID F S + + AV+V

Sbjct: 242 SYMEDLHLIIDADLEALDQVLAFAFNMNRDTFLGNVTVIDGF-----ASTGLEAVLV 295

Query: 300 DSEWFMIIYDKLYKTTSLYNPEGLYWNYLHHHQLYSTSQFGNAVAFVKSATKPVTKVAF 359
 D +WFM+YD L+K ++ NP GLYWNY+ H Q S S+F NAVAFV VT+V +

Sbjct: 296 DKDWFPMVYDNLHKMETVRNPRGLYWNYYHVWQTLVSRSFANAVAFVSGDVPVAVTQVIVS 355

Query: 360 SATTSVVKSSKDIALTFTPEATNQQGEVSSAPALVKATVKQTAGKATAVTVEGLEV 419

336

+V +G + V ATN + V V G +T + G
 Sbjct: 356 PNIAAVKQGGQQFT---AYVRATNAKDHKV-----VWSVEGGSTGTAI----TG 398

Query: 420 QSLVTFTAIGGQATVLVTV 439

L++ + Q TV TV

Sbjct: 399 DGLLSVSGNEDNQLTVKATV 418

Query= sid|110160|lan|182ORF005 Phage 182 ORF|12651-13700|3
 (349 letters)

>gi|137932|sp|P15132|VG13_BPPH2 MORPHOGENESIS PROTEIN 1 (LATE
 PROTEIN GP13) >gi|75858|pir||WMBP23 gene 13 protein -
 phage phi-29 >gi|215331 (M14782) morphogenesis protein
 [Bacteriophage phi-29] >gi|225368|prf||1301270H gene 13
 [Bacteriophage phi-29]
 Length = 365

Score = 51.5 bits (121), Expect = 8e-06
 Identities = 44/166 (26%), Positives = 70/166 (41%), Gaps = 14/166 (8%)

Query: 6 NEQIARGQTIAKILSKYGYNKSQVGVVANLHWESA---GLNPNSNEXXXXXXXX-QWT 61
 +E Q I LS G+ K + G++ N+ ES GL N +E QWT
 Sbjct: 12 SEMKVAQYILNYLSSNGWTKQAICGMLGNMQSESTINPGLWQNLDEGNTSLGFGLVQWT 71

Query: 62 PKSNLYRQAQICGLSNAKAETLEGQAEIIAQGDKTGQWMDNTPVSSAGYTNPQTLSAFKQ 121
 P SN A GL ++ II + + QW++ ++ Y K
 Sbjct: 72 PASNYINWANSQGLPYKMDMS--ELKRIIWEVNNAQWINLRDMTFKEY-----IKS 121

Query: 122 SANIDVATINFMCHWERPGKLHIEERLDAQAYSKHIDSGGGGGVK 167
 + + F+ +ERP + ER D A+ + K++ G GGGG++
 Sbjct: 122 TKTRELAMIFLASIERPANPNQPERGDQAEYWKYKNSLGGGGGGLQ 167

Query= sid|110161|lan|182ORF006 Phage 182 ORF|14995-16026|1
 (343 letters)

>gi|137945|sp|P07541|VG16_BPPZA ENCAPSIDATION PROTEIN (LATE PROTEIN
 GP16) >gi|75861|pir||WMBP16 gene 16 protein - phage PZA
 >gi|216065 (M11813) morphogenesis protein C
 [Bacteriophage PZA]
 Length = 332

Score = 402 bits (1023), Expect = e-111
 Identities = 186/332 (56%), Positives = 244/332 (73%), Gaps = 2/332 (0%)

Query: 11 EKNLYYNPNNALGFNCMLFVIGARGIGKTYGYKKFVVNRFIKHGEQFIYLRPFKTELKK 70
 +K+L+YNP L ++ ++ FVIGARGIGK+Y K + +NRFIK+GEQFIY+RR+K EL K
 Sbjct: 2 DKSIFYNPQKMSYDRILNLFVIGARGIGKSYAMKVYPINRFIKYGEQFIYVRRYKPELAK 61

Query: 71 IPQFFKTMKEFPDHLKLEVKGEFYCDKLMGWAVPLSTWGIEKSNEYPEVRTILFDEF 130
 + +F +A+EFPDH+L VKG+ FY D KL GWA+PLS W EKS N YP V TI+FDEF+
 Sbjct: 62 VSNFYNDVAQEFDPDHLVVKGRFYIDGKLAWAIPLSVWQSEKSNAYPNVSTIVFDEFI 121

Query: 131 IEKSKITYLPNEAEALLNMMETVFRRTNTRCVMLSNATSVVNPFYFLYNLQPDNLNKR 190
 EK Y+PNE ALLN+M+TVFR R RC+ LSNA SVVNPFYFL+FNL PD+NKR 190
 Sbjct: 122 REKDNSNYIPNEVSALLNMDTVFRNRERVRCICLSNAVSVVNPFYFLFFNLVDPVNKR 181

Query: 191 LYQDRGILIELCDKDFAEVKRETPFGRLIRGTEYEDFSINNEFVNDSDTFIEKRSKNSS 250
 +Y D LIE+ DS DF+ +R+T FGRLI GTEY + S++N+F+ DS FIEKRSK+S
 Sbjct: 182 VYDD--ALIEIPDSLDFSSERRKTRFGRLIDGTEYGEMSLDNQFIGDSHVFIKRSKDSK 239

Query: 251 FLCAIAFEGKIFGYWIDAETGCVVSYDYQPNTNHFYAMTTKDHEENRLLMKGNWNNYYL 310
 F+ +I + G G W+D G +YV + P+T + Y +TT D EN +L+ N++NNY+L
 Sbjct: 240 FVFSIVYNGFTLGVWVDVNGQLMYVDTAHPSTKNVYTLTTDDLNNMMLITNYKNYYHL 299

Query: 311 STVAKAFKNSYLRFDNIVIKNLHYDLFNKMKI 342
 +A AF N YLRFDN VI+N+ Y+LF KM+I
 Sbjct: 300 RKLASAFMNGYLRFDNQVIRNIAVELFRKMRI 331

Query= sid|110162|lan|182ORF007 Phage 182 ORF|7795-8775|1
 (326 letters)

>gi|1429239|emb|CAA67658| (X99260) upper collar protein
 [Bacteriophage B103]

337

Length = 308

Score = 271 bits (685), Expect = 6e-72
 Identities = 131/275 (47%), Positives = 187/275 (67%), Gaps = 5/275 (1%)

Query: 36 YYEHYRRQLTLLTFQLFEWENLPKSIDPRYLEIALHTNGYLGFFKDPTLGFMVCAGAEDG 95
 +Y HY + L L +QLFEWE LP S+DP YLE ++H GY+GF+KDP +G++ C GA G
 Sbjct: 22 WYHYHYQYLCSLAYQLFEWERLPPSVDPSPYLEXSIHQFGYVGFYKDPRIGYIACQGALSG 81

Query: 96 QIDHYHNPIFFTANEAMYHKRYPVLRYYDDDDKSKCIMLYNNDLKVPRTLPSLHRFALDMA 155
 +DHY+ P F A+ Y + + Y D +K+ + +YNNDLK TLP+L FA D+A
 Sbjct: 82 TVDHYNLPDRFHASSVGQNTFKLYNYSMDKEKNMGVAIYNNDLKSTLPALEMFAQDLA 141

Query: 156 DINQISRVNRRRAQKTPVLIQTDKQYFSLQAYNQIDENNQAVFVDKMEFDESFNWVQT 215
 ++ +I VN+ AQKTPV+I ++ SL YNQ + N +FV + ++ D + V++T
 Sbjct: 142 ELKEIIAVNQNAQKTPVLIAANDNNQLSLKNIYNQYEGNAPVIFVHESLDLD-NLKVFKT 200

Query: 216 NAPPYVVDKLRSELNEVWNEVLTFLGINNANVDKTARVQTSEVLSNNEQIESSGNILLKSR 275
 +APYVVDKL ++ N VVNEV+T+LGI NAN++K R+ TSEV SN+EQIESSGNI LK+R
 Sbjct: 201 DAPPYVVDKLNQAQNAVWNEVMTYLGIKNANLEKKERMVTSEVDSNDEQIESSGNIYLKAR 260

Query: 276 KEFCDRVNRVFGDELGDGKIDVKFRITDAVRQLQALAA 310
 +E C++++ ++G L VKFR D V Q++L A
 Sbjct: 261 QEACNKISELYGLNL----KVKFRYDIVEQMRLLA 291

Query= sid|110163|lan|182ORF008 Phage 182 ORF|14105-14983|2
 (292 letters)

>gi|4210750|emb|CAA10710| (AJ132604) LysL protein [Lactococcus
 lactis]
 Length = 235

Score = 139 bits (347), Expect = 2e-32
 Identities = 85/210 (40%), Positives = 114/210 (53%), Gaps = 14/210 (6%)

Query: 2 MNGIDISSYQTGIDLSKVPDFVNIKATGGTGYVNPDCDRAQQALSGLKKIGVYHFAHE 61
 MNGIDISSYQ ++ VP DFV IKAT GT Y+NP + Q + K +G YHFA
 Sbjct: 1 MNGIDISSQAELNAGIVPSDFVVIKATEGTNYINPTWEEQAGQVIQTNKLLGFYHFAS- 59

Query: 62 RGLEGTPOQEAQFFLDNIKG YIGKAVLILDFEGS--NQKDVNWAKAFLDYVYNKTGVKAW 119
 G P EA FF+ +K YIGKAVL+LDPE N A+ FL+ V KTG+
 Sbjct: 60 ---VGNPIAEADFFISVVKNYIGKAVLVLDPEAGAINAWGNVGARQFLNRVKEKTGINPM 116

Query: 120 FYTYTANLNTTDFSSIAKGDYGLWVAEYGSNQPOGYSQPAPPKTNN-----FPVACQF 174
 Y + ++S+I+ + LWVA+Y S P GY + P T+ + A Q+
 Sbjct: 117 IYMSSDVTRQFNWSTISSTN-PLWVAQYASMNPTGYQ--SEPWTGKGYGAWSSAAIHQY 173

Query: 175 TSKGRLPGYNGNLDLNVFYGDGNTWDLYVG 204
 +S G L ++GNLD+N+ Y + N W G
 Sbjct: 174 SSAGSLSNWSGNLDINLAYINANQWKS LAG 203

Query= sid|110164|lan|182ORF009 Phage 182 ORF|8765-9601|2
 (278 letters)

>gi|1429240|emb|CAA67659| (X99260) lower collar protein
 [Bacteriophage B103]
 Length = 293

Score = 180 bits (451), Expect = 1e-44
 Identities = 115/296 (38%), Positives = 161/296 (53%), Gaps = 33/296 (11%)

Query: 3 LKRYIESFTYYQPELS RERIEVGRKQLDFDYPFYDETKRAEFETKFINHYFLREIGSE 62
 L YIE ++ Y+ LS E+IE GR +LDFD YP +DE+ R FET FI +FY+REIG E
 Sbjct: 8 LSTYIEMWSQYETGLSMAEKIEKGRPKLDFDQYPIFDES YRKVFETHFIRNFYMRIGFE 67

Query: 63 TMGSPKFNLD EYLNLMNPYWNKMFSLNLEEF-PIFDDMDYTIDEKQKLLNEIDTNIKANR 121
 T G KFNFL+ +L +NMPY+NK+F S L ++ P+ + T K+ DT NR
 Sbjct: 68 TEGLFKFNLETWLIINMPYFNKLFES ELIKYDPLENTRLNNTGNKKN-----DTERNDNR 122

Query: 122 D-----ESKNQTKQVDQTDNRNKNTRDTGTT-----DSFSRNTYTDTPQKDLRIASNG 169
 D + K+ TK D+T+ + D TT D+F+R +D P L + +N

338

Sbjct: 123 DTTGSMKADGKSNTKTSKDTNATGSSKEDGKTGVSITDDNFNRKIDSDQPD SRLNLTTN- 181

Query: 170 DGTGVINYATNITEDLSKETTSSTGVETNNDKTNQNTSRNAS-----EKETKNTD 219
 DG G + YA+ I E+ + ++TG TNN ++ + S S T N

Sbjct: 182 DGQGTLEYASAEENNTNKRNTTG--TNNVTSSAESESTGSGTSDTVTTDNANTTTNDK 239

Query: 220 INKQONQTKDTITRYKGGKNGTDYADLLEKYRRSVLRIEKMIFREMKEGLFLLVY 275
 +N N +D I GK G YA L++ YR ++LRIEK IF EM + LF+LVY

Sbjct: 240 LNSQINNVEDYIESKIGSGTQSYASLVQDYRAALLRIEKRFDEMGE--LFMLVY 293

Query= sid|110165|lan|182ORF010 Phage 182 ORF|1310-2155|2
 (281 letters)

>gi|135604|sp|P06812|TERM BPNF DNA TERMINAL PROTEIN
 >gi|75815|pir|ERBPNP terminal protein - phage NF
 >gi|579177|emb|CAA68440| (Y00363) gene E product (AA
 1-267) [Bacteriophage NF]
 Length = 266

Score = 74.9 bits (181), Expect = 6e-13
 Identities = 73/275 (26%), Positives = 129/275 (46%), Gaps = 37/275 (13%)

Query: 3 VRISKNDRAKLEKIYGKSNKARKKYNRLRQK-GVE---ERQLPTVPTSKKRLIDYVKSTN 58
 +RI+ ND+A K+ K+ KA K +R ++K G++ E +LP + + +

Sbjct: 7 IRITNNDKALYAKLV-KNTKA--KISRTKKYIGIDLSNEIELPPLESFQ----- 52

Query: 59 MSRSDFNKMLDELVDFAQPYNENYIFEINKRNVAISRQIKEAQIKTEQAQKAKEEHYKE 118
 +R +FNK + F N+NY F NK + S+A+I E T++AQ+ +E +E

Sbjct: 53 -TREFNKKWKQKQESFTNRANQNYQFVKNKYGIVASKAKINEIAKNTKEAQRIVDEQREE 111

Query: 119 L-----NKVEVKKPTENTIVTPTILTELGADLPFQAIPDFNIDAFTSPEGVQSYLEN 170
 + K + I++P+ +T G P DFN D S +++ E

Sbjct: 112 IEDKPFISGGKQGTGQRMQILSPSQVT--GISRP---SDFNFDDVRSYARLRTLEEG 165

Query: 171 IG-KQDEQYFDERDQLYYDNFRQAMFTIFNSD--ADDIVRLDLSMGLDLFMKTYVSNFLD 227
 + K Y+D R + NP + + FNSD +D++V L + D F + Y+ F +

Sbjct: 166 MAEKASPDYDRMTQMHNQNFIEIVEKSFNSDWLSDELVERLKKIPDDFFELYLM-FDE 224

Query: 228 MNLDYIYDEAEVQKKEQVYSKIAKVIESETGGEV 262
 ++ +Y E E + E + +KI ++ G+V

Sbjct: 225 ISFEYFDSEGEDVEASEAMLNKIHSYLDYERGDV 259

Query= sid|110166|lan|182ORF011 Phage 182 ORF|9607-10158|1
 (183 letters)

>gi|1429241|emb|CAA67660| (X99260) pre-neck appendage protein
 [Bacteriophage B103]
 Length = 860

Score = 50.8 bits (119), Expect = 6e-06
 Identities = 29/105 (27%), Positives = 56/105 (52%), Gaps = 6/105 (5%)

Query: 8 KRFDGLPAVFKERFSKYPHTEYRYELLLDEEVSAIAYLNEVGALVNDMSGYLYNFIEHF 67
 +RF+ L + + + +Y T + + L E+++ +I YLN++G L ND+ N +E

Sbjct: 7 RRFEKLGEMMVQVYERYLPTAFDESMTLLEKMKIIEYLNQIGRLTNDVVEEWNKVMWEI 66

Query: 68 V-EKLEBITNDTLKKWLSDGTLENLINDTVFANYIKEIKRLQILV 111
 + + LE+ +TL+KW +G +L+ I E+K+ + V

Sbjct: 67 LNDGLEDYVKETLEKWEYEGKFADLV-----IQVIDELKQFGVSV 106

Query= sid|110169|lan|182ORF014 Phage 182 ORF|13716-14108|3
 (130 letters)

>gi|137936|sp|P11188|VG14 BPPH2 LYSIS PROTEIN (LATE PROTEIN GP14)
 >gi|75860|pir|WMBP29 gene 14 protein - phage phi-29
 >gi|15678|emb|CAA28631| (X04962) gene 14 product (AA

339

1-393) [Bacteriophage phi-29] >gi|225369|prf||1301270J
 gene 14 [Bacteriophage phi-29]
 Length = 131

Score = 96.7 bits (237), Expect = 6e-20
 Identities = 53/131 (40%), Positives = 81/131 (61%), Gaps = 3/131 (2%)

Query: 1 MIEYITQWL-ADDNHLVYGLIWLVMAMIIDFVLGFTIAKFNKEIDFSSFKAKAGIIVKV 59
 MI ++ +L D+ L+Y L +LMV M++D VLG AK N I FSSFK K G+++KV
 Sbjct: 3 MIAWMQHPLETDETKLIYWL-T-FLMVCMVVDTVLGVLFKLNPNIKFSSFKIKTGVLIVK 61

Query: 60 AEMVLVVYFIPVAVKFGAVGITMYITMLVGLILSEIYSILGHISDIDDDNNWTDYVKKFL 119
 +EM+L + IP AV F A G+ + T+ L +SEIYSI GH+ +DD +++ + ++ F
 Sbjct: 62 SEMILALAIIPFAVPPFA-GLPLLYTVYTALCVSEIYSIFGHLRLVDDKSDFLLEILENFF 120

Query: 120 DGTLNKDDIK 130
 T + + K
 Sbjct: 121 KRTSGKNKEEK 131

Query= sid|110170|lan|182ORF015 Phage 182 ORF|854-1225|2
 (123 letters)

>gi|15670|emb|CAA24483| (V01155) reading frame 10 (may be gene 4)
 [Bacteriophage phi-29]
 Length = 124

Score = 69.9 bits (168), Expect = 6e-12
 Identities = 39/119 (32%), Positives = 64/119 (53%), Gaps = 3/119 (2%)

Query: 3 IVKSTFDQTPEGMLQVFNATNGASIPLRNAI-GEVLELKDILVYSDEVSGFGGAEPSSQA 61
 IVK+TFDT+T EG +++FNA G +N G ++E I Y +G A+ +
 Sbjct: 6 IVKATFDTTETLEGQIKIFNAQTGGGQSFKNLPDGTIIEANAIAQYKQVSDTYGDAK--EE 63

Query: 62 ELVAFFTEDGKTYAGVSAVATKSAKNLIDMMTANPDIKPKISFVEGKSNGGQKFVNLOV 120
 + F DG Y+ +S ++A +LID++T + K+ V+G S+ G F +LQ+
 Sbjct: 64 TVTTIFAADGSLYSAISKTVAEASDLIDLVTTRHKLETFKVKVQGTSSKGNVFFSLQL 122

Query= sid|110174|lan|182ORF019 Phage 182 ORF|4323-4613|3
 (96 letters)

>gi|1429235|emb|CAA67654| (X99260) head morphogenesis protein
 [Bacteriophage B103]
 Length = 101

Score = 60.9 bits (145), Expect = 1e-09
 Identities = 34/96 (35%), Positives = 53/96 (54%), Gaps = 5/96 (5%)

Query: 1 MEIKEHESILNGILESVDGEARSKIVEHLEALREDYGATTEALTSANSTLEKLKKNED 60
 ME HE ILN + + + R+++ L+ LR DY+ + S EKL+ +N
 Sbjct: 3 MERDSHEEILNKLNDPELEHSERTEL---LQQLRADYGSVLSEFSELTATEKLRAENS 59

Query: 61 LVISNSKLFRRERAIVEPAEN--NEPETDQNITLDDL 94
 L++SNSKLF+ I + E + E + IT++DL
 Sbjct: 60 LIVSNSKLFQVGITKEKEEIKQEELSETITIEDL 95

Query= sid|110180|lan|182ORF025 Phage 182 ORF|548-814|2
 (88 letters)

>gi|138099|sp|P06955|VG6_BPPZA EARLY PROTEIN GP6
 >gi|75841|pir|ERBP62 gene 6 protein - phage PZA
 >gi|216047 (M11813) gene 6 product [Bacteriophage PZA]
 >gi|224746|prf||1112171K ORF 6 [Bacteriophage PZA]
 Length = 96

Score = 55.0 bits (130), Expect = 8e-08
 Identities = 28/79 (35%), Positives = 45/79 (56%)

340

Query: 4 KLMQRNVTSTKVEFSEVIVQDGAPTIVPCEPVVLTGKLSEEKALSAIKRINPDINVVVTN 63
K+MQR +T T V +++++ DG + G LS E+A +KRK + V V +
Sbjct: 3 KMMQREITKTTVNVAKMVMVDGEVQVEQLPSETFVGNLSMEQAQWRMKRKYKGEVPVQVVS 62

Query: 64 VSHETALYTMPVDKFIELD 82
V T +Y +PV+KF+E+A
Sbjct: 63 VEPNTEVYELPVEKFLEVA 81

Table 26

Secondary structure prediction for ORF 182ORF008

```

1  MMNGIDISSY QTGIDLSKVP CDFVNIKATG GTGYVNPDCD RAFQQALSLG KKIGVYHFAH
   CCCCCCCCCC CCCCCCCCCC CEEEEEEEC CCCCCCCCCC HHHHHHHHHC CCCCEEEEEE
61  ERGLEGTPOQ EAQFFLDNIK GYIGKAVLIL DFEGSNQKDV NWAKAFLDYV YNKTGVKAWF
   CCCCCCCHH HHHHHHHHHC CCCCEEEEEE CCCCCCHH HHHHHHHHHC HCCCCEEEEE
121 YTYTANLNTT DFSSIAKGDY GLWVAEYGSN QPQYSQPAP PKTNNFPIVA CFQFTSKGRL
   EEECCCCCCC CCCECCCCC CEEEECCCC CCCCCCCCCC CCCCCCCEE EEECCCCCCC
181 PGYNGNLDLN VFYGDGNTWD LYVGKKQDQI VPPENKIFDA TSDEFIFTLT TGSTSVFYFD
   CCCCCCCEE EEECCCCCE EEECCCCCCC CCCCCCCCCC CCCEEEEEEC CCCCEEEEC
241 GETIFELSDP TQLDHIRGTY NHVHGKEIPS MVWTPEQFDI YLKMYEKKPV YK
   CEEEECCCC CCHHHHCCEE CCCCCCECC CCCCCCHH HHHHCCCCCE EC

```

Secondary structure prediction for ORF 182ORF014

```

1  MIEYITQWLA DDNHLVYGLI IWLVMAMIID FVLGFTIAKF NKEIDFSSFK AKAGIIVKVA
   CCCCECCCC CCCCHHHHH HHHHHHHHHC CCCCCHHHH HHCCEEEEEE
61  EMVLVVFIP VAVKFGAVGI TMYITMLVGL ILSEIYSILG HISDIDDDNN WTDYVKKFLD
   EEEEEEECC CEECCCEE EEEEEEEEE EEEEEEECC CCCCCCCCC CEEEEEECC
121 GTLNRKDDIK
   CCCCCCEEC

```

Table 27

Enterococcus accession numbers 242/242

gi 2895751 gb AF044978.1 AF044978 [2895751]	gi 4098267 gb U76614.1 BLU76614 [4098267]
gi 4803755 dbj AB026843.1 AB026843 [4803755]	gi 47019 emb Y00116.1 SFAMB1 [47019]
gi 4769001 gb AF140549.1 AF140549 [4769001]	gi 4158179 emb AL035206.1 SC9B5 [4158179]
gi 4760901 gb AF099088.1 AF099088 [4760901]	gi 4165458 emb X79343.1 EF16SSPA [4165458]
gi 4704705 gb AF121254.1 AF121254 [4704705]	gi 4165457 emb X79342.1 EFTRNALA [4165457]
gi 3342117 gb AF076604.1 AF076604 [3342117]	gi 4165456 emb X79341.1 EF23SRNA [4165456]
gi 4688824 emb AJ132470.1 ESP132470 [4688824]	gi 4150978 emb Y14027.1 EFY14027 [4150978]
gi 4732085 gb AF125553.1 AF125553 [4732085]	gi 4127803 emb AJ223161.1 EFAJ3161 [4127803]
gi 4732082 gb AF125552.1 AF125552 [4732082]	gi 2956685 emb Y16413.1 EFENTIJO [2956685]
gi 4732079 gb AF125551.1 AF125551 [4732079]	gi 2665346 emb Y13922.1 EHY13922 [2665346]
gi 4732076 gb AF125550.1 AF125550 [4732076]	gi 4324675 gb AF109375.1 AF109375 [4324675]
gi 4732073 gb AF125548.1 AF125548 [4732073]	gi 4234627 gb AF061013.1 AF061013 [4234627]
gi 4732070 gb AF125547.1 AF125547 [4732070]	gi 4234626 gb AF061012.1 AF061012 [4234626]
gi 4732067 gb AF125546.1 AF125546 [4732067]	gi 4234625 gb AF061011.1 AF061011 [4234625]
gi 4732064 gb AF125545.1 AF125545 [4732064]	gi 4234624 gb AF061010.1 AF061010 [4234624]
gi 4732061 gb AF125544.1 AF125544 [4732061]	gi 4234623 gb AF061009.1 AF061009 [4234623]
gi 4704653 gb AF114715.1 AF114715 [4704653]	gi 4234622 gb AF061008.1 AF061008 [4234622]
gi 4704564 gb AF102550.1 AF102550 [4704564]	gi 4234621 gb AF061007.1 AF061007 [4234621]
gi 4688827 emb AJ238249.1 EFA238249 [4688827]	gi 4234620 gb AF061006.1 AF061006 [4234620]
gi 4680606 gb AF125198.1 AF125198 [4680606]	gi 4234619 gb AF061005.1 AF061005 [4234619]
gi 4633279 gb AF117609.1 AF117609 [4633279]	gi 4234618 gb AF061004.1 AF061004 [4234618]
gi 4633124 gb AF110130.1 AF110130 [4633124]	gi 4234617 gb AF061003.1 AF061003 [4234617]
gi 4590399 gb AF124258.1 AF124258 [4590399]	gi 4234616 gb AF061002.1 AF061002 [4234616]
gi 4590336 gb AF108380.1 AF108380 [4590336]	gi 4234615 gb AF061001.1 AF061001 [4234615]
gi 4590335 gb AF108379.1 AF108379 [4590335]	gi 4234614 gb AF061000.1 AF061000 [4234614]
gi 4019167 gb U21300.1 CXU21300 [4019167]	gi 3138990 gb AF060241.1 AF060241 [3138990]
gi 4545122 gb AF077816.1 AF077816 [4545122]	gi 3138986 gb AF060240.1 AF060240 [3138986]
gi 4433610 gb AF106614.1 AF106614 [4433610]	gi 4204535 gb AF094803.1 AF094803 [4204535]
gi 4468838 emb AJ132039.1 EFA132039 [4468838]	gi 4204534 gb AF094802.1 AF094802 [4204534]
gi 4468121 emb AJ132958.1 BPH132958 [4468121]	gi 4204533 gb AF094801.1 AF094801 [4204533]
gi 4456104 emb Y17302.1 EHI17302 [4456104]	gi 4204532 gb AF094800.1 AF094800 [4204532]
gi 4433611 gb AF106615.1 AF106615 [4433611]	gi 4204531 gb AF094799.1 AF094799 [4204531]
gi 4433607 gb AF106611.1 AF106611 [4433607]	gi 4204530 gb AF094798.1 AF094798 [4204530]
	gi 4204529 gb AF094797.1 AF094797 [4204529]
	gi 4204528 gb AF094796.1 AF094796 [4204528]
	gi 4204527 gb AF094795.1 AF094795 [4204527]

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 gi|4204517|gb|AF094785.1|AF094785 [4204517]
 gi|4204516|gb|AF094784.1|AF094784 [4204516]
 gi|4204515|gb|AF094783.1|AF094783 [4204515]
 gi|4204514|gb|AF094782.1|AF094782 [4204514]
 gi|4204513|gb|AF094781.1|AF094781 [4204513]
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 gi|4151367|gb|AF093508.1|AF093508 [4151367]
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 gi|2828135|gb|AF039902.1|AF039902 [2828135]
 gi|2828134|gb|AF039901.1|AF039901 [2828134]
 gi|2828133|gb|AF039900.1|AF039900 [2828133]
 gi|2828132|gb|AF039899.1|AF039899 [2828132]
 gi|2828131|gb|AF039898.1|AF039898 [2828131]
 gi|4103866|gb|AF028812.1|AF028812 [4103866]
 gi|4103864|gb|AF028811.1|AF028811 [4103864]
 gi|2605925|gb|AF029727.1|AF029727 [2605925]
 gi|1402750|gb|U60038.1|EFU60038 [1402750]
 gi|1835780|gb|U86375.1|EFU86375 [1835780]
 gi|3831555|gb|AF047608.1|AF047608 [3831555]
 gi|3790617|gb|AF097414.1|AF097414 [3790617]
 gi|3767587|dbj|AB005036.1|AB005036 [3767587]
 gi|3757810|gb|AF042288.1|AF042288 [3757810]
 gi|3747039|gb|AF093509.1|AF093509 [3747039]
 gi|3660559|dbj|AB017811.1|AB017811 [3660559]
 gi|1147743|gb|U42211.1|EHU42211 [1147743]
 gi|3676412|gb|AF051917.1|AF051917 [3676412]
 gi|3676164|emb|AJ011113.1|EFA011113 [3676164]
 gi|2612869|gb|AF005726.1|AF005726 [2612869]
 gi|2353762|gb|AF016233.1|AF016233 [2353762]
 gi|2149899|gb|U94707.1|EFU94707 [2149899]
 gi|2149149|gb|U82366.1|LSU82366 [2149149]
 gi|1469463|gb|U49512.1|EFU49512 [1469463]
 gi|1244503|gb|U35366.1|EFU35366 [1244503]
 gi|833854|gb|U26268.1|EFU26268 [833854]
 gi|841200|gb|U18931.1|CPU18931 [841200]
 gi|460079|gb|U00457.1|U00457 [460079]
 gi|460077|gb|U00456.1|U00456 [460077]
 gi|535661|gb|L34675.1|INSTRANSPO [535661]
 gi|3023041|gb|AF007787.1|AF007787 [3023041]
 gi|431124|gb|L15633.1|TRN916ENT [431124]
 gi|388106|gb|L23802.1|ENEEBSA [388106]
 gi|3608387|gb|AF071085.1|AF071085 [3608387]
 gi|3551851|gb|AF076027.1|AF076027 [3551851]
 gi|3551773|gb|U94770.1|SPU94770 [3551773]
 gi|3551743|gb|U57498.1|ECU57498 [3551743]
 gi|3243178|gb|AF063010.1|AF063010 [3243178]
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Table 28

Phage Dp1 complete genome sequence. 56506 nucleotides.

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Table 29

Phage dp1 ORFs list

nb	Name	Frame	Position	Size (a.a.)	Key words
1	dp1ORF001	2	36698..40390	1230	Putative tail;
2	dp1ORF002	1	32386..35835	1149	Tail;
3	dp1ORF003	3	53538..55877	779	DNA polymerase I;
4	dp1ORF004	3	40401..42440	679	Minor structural;
5	dp1ORF005	1	23674..25434	586	
6	dp1ORF006	2	45296..46987	563	SWI/SNF Helicase;
7	dp1ORF007	3	22230..23621	463	Terminase;
8	dp1ORF008	1	49624..50961	445	DNA Helicase;
9	dp1ORF009	2	13160..14404	414	
10	dp1ORF010	2	8699..9859	386	RecA;
11	dp1ORF011	3	28017..29096	359	Major head;
12	dp1ORF012	3	5346..6419	357	DNA pol. III beta;
13	dp1ORF013	3	10215..11240	341	DNA pol. III gamma and tau;
14	dp1ORF014	3	50961..51974	337	DNA primase;
15	dp1ORF015	1	3793..4728	311	
16	dp1ORF016	3	43413..44303	296	Amidase;
17	dp1ORF017	1	11242..12081	279	
18	dp1ORF018	3	35847..36686	279	
19	dp1ORF019	2	12161..12967	268	
20	dp1ORF020	1	1864..2658	264	exsD; Coenzyme PQQ;
21	dp1ORF021	2	2504..3295	263	GTP cyclohydrolase;
22	dp1ORF022	2	30896..31675	259	
23	dp1ORF023	2	6419..7195	258	
24	dp1ORF025	-1	18026..18778	250	
25	dp1ORF024	3	25992..26738	248	
26	dp1ORF026	2	21512..22252	246	
27	dp1ORF027	1	52762..53490	242	
28	dp1ORF028	3	44595..45299	234	
29	dp1ORF029	2	662..1348	228	exsB;
30	dp1ORF031	3	26943..27611	222	
31	dp1ORF030	-2	19423..20088	221	
32	dp1ORF032	1	52033..52647	204	
33	dp1ORF033	2	7670..8239	189	
34	dp1ORF035	-1	16859..17425	188	
35	dp1ORF036	1	48808..49362	184	DNA replication;
36	dp1ORF037	1	55855..56388	177	
37	dp1ORF034	2	131..652	173	
38	dp1ORF038	3	1350..1871	173	exsC; 6-pyruvoyltetrahydropterin;
39	dp1ORF039	3	3306..3803	165	Citrulline biosynthesis;
40	dp1ORF040	1	7192..7683	163	
41	dp1ORF041	3	8208..8699	163	dUTPase;
42	dp1ORF042	1	48082..48561	159	
43	dp1ORF043	1	31699..32154	151	
44	dp1ORF044	-1	25211..25666	151	
45	dp1ORF045	2	25340..25777	145	
46	dp1ORF046	3	42774..43202	142	
47	dp1ORF047	1	47542..47961	139	
48	dp1ORF048	-3	16308..16709	133	
49	dp1ORF049	-3	43620..44018	132	
50	dp1ORF050	3	15081..15476	131	
51	dp1ORF051	2	29765..30154	129	
52	dp1ORF053	-3	49917..50300	127	
53	dp1ORF052	3	30516..30893	125	
54	dp1ORF054	2	14423..14800	125	
55	dp1ORF055	3	27627..28004	125	
56	dp1ORF056	-3	18780..19151	123	
57	dp1ORF057	1	9859..10218	119	
58	dp1ORF058	3	15633..15989	118	
59	dp1ORF059	1	30154..30507	117	
60	dp1ORF060	-2	37717..38070	117	
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63	dp1ORF064	2	29108..29449	113	

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66	dp1ORF068	3	29451..29768	105	
67	dp1ORF069	-3	20094..20411	105	
68	dp1ORF061	-3	19161..19475	104	
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83	dp1ORF065	-3	51246..51497	83	
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85	dp1ORF087	-2	29794..30036	80	
86	dp1ORF088	3	5040..5279	79	
87	dp1ORF089	-2	12256..12495	79	
88	dp1ORF273	3	56256..56486	76	
89	dp1ORF078	-3	17280..17507	75	
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91	dp1ORF091	1	43189..43413	74	Holin;
92	dp1ORF092	3	46989..47213	74	
93	dp1ORF093	-2	45538..45756	72	
94	dp1ORF095	3	8877..9089	70	
95	dp1ORF096	-1	46469..46681	70	
96	dp1ORF097	-1	38888..39100	70	
97	dp1ORF098	1	43627..43836	69	
98	dp1ORF099	3	38298..38507	69	
99	dp1ORF100	1	1597..1803	68	
100	dp1ORF101	2	19220..19426	68	
101	dp1ORF094	1	8281..8484	67	
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103	dp1ORF104	-1	21224..21427	67	
104	dp1ORF105	-2	1828..2028	66	
105	dp1ORF106	-3	10329..10529	66	
106	dp1ORF108	-1	49250..49447	65	
107	dp1ORF109	-2	31435..31632	65	
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125	dp1ORF127	-3	13335..13511	58	
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145	dp1ORF148	-1	28484..28636	50	
146	dp1ORF150	-3	15033..15185	50	
147	dp1ORF134	-2	349..498	49	
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266	dp1ORF265	-1	4700..4801	33	
267	dp1ORF266	-2	50119..50220	33	
268	dp1ORF267	-2	47266..47367	33	
269	dp1ORF268	-2	12520..12621	33	
270	dp1ORF269	-3	53733..53834	33	
271	dp1ORF270	-3	50691..50792	33	
272	dp1ORF271	-3	19638..19739	33	
273	dp1ORF272	-3	1455..1556	33	

Table 30

Predicted Dp-1 amino acid sequences

dp1ORF001

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36698 atgattgacaataatttacctatgagtcgaattcctggcgaaattgttcaagtatatgaccaaacttcaatctaattggagca
1 M I D N N L P M S P I P G E I V Q V Y D Q N F N L I G A
36782 agtgatgaaatcttagcaagcattacgaagcgaattgtgactcgagctcgaggaaaagaacttctcatttggaaagtatt
29 S D E I F S K H Y E D E I V T R A R G K E T F T F E S I
36866 gaaacctcatctatctatcaacacttaaggttgaaacattatccagtatggaggaagatgggttccgaattaaatatgctcag
57 E T S S I Y Q H L K V E N I I Q Y G G R W F R I K Y A Q
36950 gacgtagaagatgtcaaagggcttaccaggttacctgctacgcattatgggtatgaactagcagaaggcttgcctaggaagtgtg
85 D V E D V K G L T K F T C Y A L W Y E L A E G L P R K L
37034 aaacacgttgcttcttctgtaggcgctgtgcgctagatattatcaaagacgcaggtgaatgggttgcactagtgttgcctcctt
113 K H V A S S V G A V A L D I I K D A G E W V R L V C P P
37118 gacgggtgctaaacaagaagtcgaagcataacagcgcagaaaaattcaatgctttggcatcttcgatattcttgcagaagcaatac
141 D G A N K Q V R S I T A A E N S M L W H L R Y L A K Q Y
37202 aatttgaattgacatttgggtttagaagaattatcaagcaagaggttagaattgttcaaaccgttatttcttcagccttat
169 N L E L T F G Y E E I I K Q E V R I V Q T V V F L Q P Y
37286 gtcgagttctaaagtagacttctctttagttagaagaaatttgaatatgtcactaggcaggaagattctcgaaacctgtgt
197 V E S K V D F P L V V E E N L K Y V T R Q E D S R N L C
37370 acggcttacaagttgacaggtaaaaaggaagagcagtcgaagccttttaacgtttgcttctatcaacaatggaagtgaatat
225 T A Y K L T G K K E E G S Q E P L T F A S I N N G S E Y
37454 ctcatgtatgttctgtgttactacacgccacatgaagcctcgatatattgctaaatctaaaagcgacgaacattttagaatt
253 L I D V S W F T T R H M K P R Y I A K S K S D E H F R I
37538 aaagaaaatttgatgagtgctgcgctgttcttgcacatctacagtcgcccactaatggatatgaggttcagcggctcctt
281 K E N L M S A A R A Y L D I Y S R P L I G Y E A S A V L
37622 tataacaaggttctgacttgcacatactcaactaattgtcgacgaccattatgatgttatcgagtgccgaaagattatctgct
309 Y N K V P D L H H T Q L I V D D H Y D V I E W R K I S A
37706 cgaaaaattgactacgacgacctttcaactctactatcttcccaagaccctcgaaaagacttgatggacttgccttaagag
337 R K I D Y D D L S N S T I I F Q D P R K D L M D L N E
37790 gacggcgaaggagtcctttcaggggaaactgtaaatgagtcaccaagttgttattagatagcagatgacattttagggactaat
365 D G E G V L S G E T V N E S Q V V I R Y A D D I L G T N
37874 tttaatgcagaatctgggaaatacattgggtgtccttaataataaagaaccgagcgaattagttcctgacgactttacattgg
393 F N A E S G K Y I G V L N T N K K P S E L V P D F T W
37958 attcgactagaaggtcctaaaggtgacgcaggtttaccgggagctcctggcggtgatggagtcgacggtgtactggaaagagc
421 I R L E G P K G D A G L P G A P G R D G V D G V P G K S
38042 ggagtaggagatagcagatcacgctatcacttatcgttatcgttccggaacgcaagagcctgaaaatggatggagcgaacaa
449 G V G I A D T A I T Y A V S V S G T Q E P E N G W S E Q
38126 gttcctgaactcataaaaggtcgattcttctgtggactcaaaacttttggagatatactgacggctcactgaaactggactactcc
477 V P E L I K G R F L W T K T F W R Y T D G S H E T G Y S
38210 gttgcttatatagggaagcggaaattccggaaaagcggaaatcgaggttaaggacggagtaggtatagccgcaactgaagtc
505 V A Y I G Q D G N S G K D G I A G K D G V G I A A T E V
38294 atgtatgcaagttcgccatctgctactgaagctccaggtgggtggatgctacgcaagttcctaccgtccaggtgggtcagtat
533 M Y A S S P S A T E A P A G G W S T Q V P T V P G G Q Y
38378 ttatggactcgaacaagatggcgctacactgaccaaactgatgaaattggatatcagtttcaagaatgggagcaggggtcctt
561 L W T R T R W R Y T D Q T D E I G Y S V S R M G E Q G P
38462 aaaggtgacgcaggtcgtgacgggtattgcaggaagacgaatagggttgaagtcacactcagttcttatggaattagtcctc
589 K G D A G R D G I A G K N G I G L K S T S V S Y G I S P
38546 actgattctgcgattctctggagtagggcttcacaagttccttcttaatacaaggtcaatatcttggactcgaactatttgg
617 T D S A I P G V W A S Q V P S L I K G Q Y L T R T I W
38630 acctataccgattcaactaccgaacgggctatcaaaaaacctacattccaaaagacgggaatgacgggtaaaaatggaattgct
645 T Y T D S T T E T G Y Q K T Y I P K D G N D G K N G I A
38714 ggtaaggatggggtaggaaattaagtcacgaccattacctacgcaggtcacaactcaggaacagttgcgctacttcaaatgg
673 G K D G V G I K S T T I T Y A G S T S G T V A P T S N W
38798 acttctgctattccaaatgttcaaccgggattcttctgtggacgaaaactgttggaaactatactgatgacactagcgaacaa
701 T S A I P N V Q P G F F L W T K T V W N Y T D D T S E T
38882 ggttactcagtttccaagatagggtgaaacaggtcctagwaggttcaaggtcctcaaggtcctcaagggttcaaggaaattcct
729 G Y S V S K I G E T G P R G V Q G L Q G P Q G L Q G I P
38966 ggacctgcaggagctgacggacgttcgcaatatctacacctcgcttctctaatagtcctaaacgggtgagggatttagctacact
757 G P A G A D G R S Q Y T H L A F S N S P N G F S H T
39050 gacagcggacgagcagcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtcagtc
785 D S G R A Y V G Q Y Q D F N P V H S K D P A A Y T W T K
39134 tggaaggggaatgacggagctcaagggataccgggaagcagcagcagcagcagcagcagcagcagcagcagcagcagcagcagc
813 W K G N D G A Q G I P G K P G A D G K T N Y F H I A Y A
39218 tcaagtgacagcaggtcacgtgagttcagttgaaagataatacaacaatatatgggttattactccgattatgagcaagfca
841 S S A D G S R E F S L E D N N Q Q Y M G Y Y S A
39302 gatagcaggatcgaactaagtatcgatgggttgaccgccttgccaatgttcaagtgaggagtcgaacacaggttcttatttct
869 D S R D R T K Y R W F D R L A N V Q V G G R N E F L N S
39386 ttatttgaatttgggtttaaaccctcgctattctagttacaaatctaattggacggacaaagatcaaacgcaaggacagatatctgct
897 L F F G L K P R Y S S Y A N L M D G Q D Q T Q Q I S A
39470 actattgacgaacgtcaacgggttcaaggtgcatactctttagacttgactcaacatggaacgggttaaacggcagaacccaaaaa
925 T I D E R Q R F K G A N S L R L D S T W N G K P Q N Q K

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364

39554 ctgaccttttcttaggaggagatagcgattaggtactccaaccgagtggtcctaatttagaaggctgatcagtttctgggct
953 L T F S L G G D T R L G T P T E W S N L E G R I S F W A
39638 aaggcctcttaggaacggagtgagcttagctgcacggcggttatcgtagtaacgtattaccgcaaccttaaccagctcaatgg
981 K A S R N G V S L A A R P G Y R S N V F T A T L T D Q W
39722 aagttctacgatttttaattctttgacaaagtttaattcaattgtaccgctgaagcaattttccatgtattcactcaaagtgt
1009 K F Y D F K F F D K V N S N C T A E A I F H V F T Q S C
39806 tcagtggtgctcaatcatattaaaatcgaacttggaatattcttactccttttagtgaagcagaggaacaccttaaatatcga
1037 S V W L N H I K I E L G N I S T P F S E A E D L K Y R
39890 attgactcaaaagccgatcaaaagctaactaaccaacagtgtagcggtcactcaggaagggctcaactacatgacgcagaactg
1065 I D S K A D Q K L T N Q Q L T A L T E K A Q L H D A E L
39974 aaagctaaggctacaatggagcagtttaagtaacttagaaaaggcttatgaaggtagaataaagctaataagaagcctatcaaa
1093 K A K A T M E Q L S N L E K A Y E G R M K A N E E A I K
40058 aaatcggaagccgactaatcttagcggaagtcgaattgaagctactatccaagaacttggtggggtacgggaactgaagaag
1121 K S E A D L I L A A S R I E A T I Q E L G L R E L K K
40142 ttctgtagcagttacatgagctcttctaatgaaggcttaattatcggttaagaacgacggtagctctaccattaaggatcaagt
1149 F V D S Y M S S S N E G L I I G K N D G S S T I K V S S
40226 gaccgaatttctatgttcttcgaggaatgaagttatgtacctacgcaagggttcattcacatcgataacgggattctttacc
1177 D R I S M F S A G N E V M Y L T Q G F I H I D N G I F T
40310 caatccattcaagtggcggttagaacggaacaataactcggttaattccagacatgaacgtgattcggtatgtaggataa
40390
1205 Q S I Q V G R F R T E Q Y S F N P D M N V I R Y V G *
dp1ORF002
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29 Q R L A L E S S K S F Q I G S A L T G L G K G L T T A V
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57 T L P L M G F A A A S I K V G N E F Q A Q M S R V Q A I
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32722 cgggctcaaggtatggaaaatctagcttcagcgggttccaggttaaatgaaatcatggagcgtatgcgggggttactgacgtg
113 A A Q G M E N L A S A G F Q V N E I M D A M P G V L D L
32806 gctgcggtatctggaggagatgtggccgcgagctccgagggccatgggtgatttcacttcgagcctttgggattagaggcaaaacag
141 A A V S G G D V A A S S E A M A S S L R A F G L E N Q
32890 ggggtcacgtgggtgacgtatttgcgcgagcagctgatacgaacgcagaaactagcgacatggcagaggcgatgaatac
169 A G H V A D V F A R A A A D T N A E T S D M A E A M K Y
32974 gtgcacccgttgctcactctatgggcttgagccttgagaaacgggtgctctattgggattatggcgcagccgggtattaaag
197 V A P V A H S M G L S L E E T A A S I G I M A D A I K
33058 ggctcgcaagccgggaaccacgcttagagcgctctctcgcgtattgccaacctacgaaagcgatggtcaaatcaatgcaggaa
225 G S Q A G T T L R G A L S R I A K P T K A M V K S M Q E
33142 ttaggagtttcggtctacgacgcgaacggaacatgattccactaagagaacaaatcgctcaactgaaaacagctactgcagga
253 L G V S F Y D A N G N M I P L R E Q I A Q L K T A T A G
33226 ctaacacaaggaagcaaatcgctcaccttggttaccttgattggccaaaactcggtgtcaggtatgcttgcaactattagacga
281 L T Q E E R N R H L V T L Y G Q N S L S G M L A L D A
33310 ggtcctgagaaattggataagatgaccaatgctctcgtgaactcggacggagctgctaaggaatggcagaaactatgcaggac
309 G P E K L D K M T N A L V N S D G A A K E M A E T M Q D
33394 aaacttgctagtaaaatcgagcaaatgggagagcttctcaggtctggtgctattattggtcaacaaactccttgagcctgcaact
337 N L A S K I E Q M G G A F E S V A I I V Q Q I L E P A L
33478 gctaaaatcggtggagcaatcacaaggtctcgaagcattcgtaaatatgtcacctatcggtcaaaagatgggtgtcatattc
365 A K I V G A I T K V L E A F V N M S P I G Q K M V V I F
33562 gcaggaaatgggttcagcccttgagccactgctcctaattgcaggaatgggtgatgacaactattgtcaagtaagaaatgctatt
393 A G M V A A L G P L L L I A G M V M T T I V K L R I A I
33646 cagtttttaggtccagcatttatgggaacgatgggaaccattgcaggagttatagcaatattctatgctctggtcgccgtgttc
421 Q F L G P A F M G T M G T I A G V I A I F Y A I V F
33730 atgatagcctacacaaaatcgagagatttagaaactttatcaacagctcttgccctgctattaaagctgggtttggaggagcg
449 M I A Y T K S E R F R N F I N S L A P A I K A G F G G A
33814 ttggaatggctacttccacgactgaaagagtttagggagaatgggtacagaaggcaggcgagaaggcgaaagctcggtcagtt
477 L E W L L P R L K E L G E W L Q K A G E K A K E F G Q S
33898 gtagggtctaaagtgtcaaaactgctcgaacagtttggaataagtatcggtcaggcaggaggtcgattggtcagttcattgga
505 V G S K V S K L L E Q F G I S I G Q A G G S I G F I G
33982 aatgttctcgaaaggctaggagggcatttggaagagtaggaggtcatttcaattgctgtttcacttgaacaaaatcggt
533 N V L E R L G G A F G K V G G V I S I A V S L V T K F G
34066 ctgcatttctagggttacaggaccactcgggattgcttagtctggttagtttcatctttgacagcttgggctagaaacaggt
561 L A P L G I T G P L G I A I S L L V S F L T A W A R T G
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617 L P V F V E K G T Q I L V K I I E G I A S A V P Q V V E
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34402 gaagcgcttataaatggctctgttcaatctcttctactatcattcaagcagctgttcaaatatcactgcttattcaatgggt
673 E A L I N G L V Q S L P T I I Q A A V Q I I T A L F N G
34486 cttgttcaggcacttctacgcttattcaagcaggtcttcaaatgttgcagctctcataaacggactagttcaagcgcttcg
701 L V Q A L P T L I Q A G L Q I L S A L I N G L V Q A L P
34570 gcaattattcaagcagctgttcaaatatcatgtcgctgttcaagcactaattgaaaacttgcttatgataatcgaagcagcg
729 A I I Q A A V Q I I M S L V Q A L I E N L P M I I E A A

365

34654 atgcagattataatgggtctagtcacgcactgattgaaaatataaggacctatcttagaagcagggtatcaaatcttaattggct
757 M Q I I M G L V N A L I E N I G P I L E A G I Q I L M A
34738 ttaatcgagggtacttattcaagtgtctcctgaactattacagcagcgattcaaatcattacttcactattagaagcaatcttg
785 L I E G L I Q V L P E L I T A A I Q I I T S L L E A I L
34822 tcgaacctctcctcaacttctagaagccggaggttaattgtcttttactcacttctcgaagggttgctaaatatgcttctcctaacta
813 S N L P Q L L E A G V K L L L S L L Q G L L N M L P Q L
34906 attgcaggggtcttgcataatcatgatggcacttcttaagcagttatcgacttcgctccctaaacttcttcaagcaggtgttcaa
841 I A G A L Q I M M A L L K A V I D F V P K L L Q A G V Q
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869 L L K A L I Q G I A S L L G S L L S T A G N M L S S L V
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897 S K I A S F V G Q M V S G G A N L I R N F I S G I G S M
35158 attggttcagctgtctctaaaattggcagcatgggaacctcaattgtttctaagggtactggattcgctggacaaatggtaagc
925 I G S A V S K I G S M G T S I V S K V T G F A G Q M V S
35242 gcaggggtcaaccttctgaggttattcaatgtatcagttccatggtaagttctcggttaagttcggtgaggtggttaattgggt
953 A G V N L V R G F I N G I S S M V S S A V S A A A N M A
35326 agcattgcattaaatgccgttaagggtattcttaggtattcacttctccttcacgtgtcatggagcagatgggtattctatacgggt
981 S S A L N A V K G F L G I H S P S R V M E Q G I Y T G
35410 caagggttcgtaaatgggtattggttaacatgattcgaaactacacgtgacaagggttaagaaatgggtgaaactgttactgaagct
1009 Q G F V N G I G N M I R T T R D K A K E M A E T V T E A
35494 ctcagcgacgtgaagatggatattcaagaaatggagttatagaaaagggttaaatcagtttacgaaaagatgggtgaccaactt
1037 L S D V K M D I Q E N G V I E K V K S V Y E K M A D Q L
35578 cctgaaactcttcagctcctgatttcgaagatgttcgtaaaagcagcgggttcgctcgagtggaactgttcaatacaggaagt
1065 P E T L P A P D F E D V R K A A G S P R V D L F N T G S
35662 gacaacctcaaccaactcagtcacaaatcctaaaacatcaaggcgagcaaacctgttgaacattggaacaaatcgtagttcga
1093 D N P N Q P Q S Q S K N N Q G E Q T V V N I G T I V V R
35746 aacaatgacgagcttgacaaactgtcgagaggattgtataatagaagtaagaaactctatcaggggttggttaacattgtaaca
1121 N N D D V D K L S R G L Y N R S K E T L S G F G N I V T
35830 ccgtaa 35835
1149 P *

dp1ORF003

53538 atggcacaaaaaggactctttggtgcaagcctcggtctagcaagaagaacgatgctcagttacttgctcaacggaaaaacagg
1 M A Q K G L F G A K P R S S K K N D A Q L L A Q R K N R
53622 aagcctgcagttgaggttactttacatttcaggaaacgctctaaaggacgcagttgcttagagctcgtactcttcaactaggatt
29 K P A V E V T Y I S G N A L K D A V A R A R A R T R I
53706 cttggacacgttcttgatagacttgagttaatcactgaggaagcaaaactcgagcagtatgtagacaaaatgattgaagacgga
57 L G H V L D R L E L I T E E A K L E Q Y V D K M I E D G
53790 atagttcttattgacgtagaaaactgatggactcattcagcatgagctggcaggtgctgcttactcactcactgcaaa
85 I G S I D V E T D G L D T I H D E L A G V C L Y S P S Q
53874 aaaggaatctatgctcctgtcaatcatgttagcaaatgacgaagatgcgaattgaagaatcaaaacttctcctcgagttcgatgaag
113 K G I Y A P V N H V S N M T K M R I K N Q I S P E F M K
53958 aaaaactttcaacggattgttagattcaggaattcctgtcatctatcataattcgaaatttgacatgaaatcgatttattggcga
141 K M L Q R I V D S G I P V I Y H N S K F D M K S I Y W R
54042 ctcggcgtcaaaatgaatgagccagcgtgggatacatatttagccgcaatgcttttaaatgaaaaacagagctccacagcttgaaa
169 L G V K M N E P A W D T Y L A A M L L N E N S H S L K
54126 agtcttactctaaatattgtaggaacgaagaaacgcagaggttgcaaaatttaattgacttatttaagggaattccttttagt
197 S L H S K Y V R N E E N A E V A K F N D L F K G I P F S
54210 ttaattcctcctgattgttccttatgtatgcgcgcctatgcgaactttcgaaactttcgaaacttctgaacaaatcgaacaaatc
225 L I P P D V A Y M Y A A Y D P L Q T F E L Y E F Q E Q Y
54294 ttgactccaggaaactgaacaaatgaagaatataacttggaagaaagttcctcatggttcttcataatattgagatgcctctaatt
253 L T P G T E Q C E E Y N L E K V S W V L H N I E M P L I
54378 aaagttctcttcgacatggaagtctacggtgtcgacttagaccaagataagctggcagaaattagagaacagtttactgccaat
281 K V L F D M E V Y G V D L D Q D K L A E I R E Q F T A N
54462 atgaacagggctgagcaagagtttcaacagcttgcagcgaatggcagcctgaaattgaagaacttcgacaaacttaattccag
309 M N E A E Q E F Q L V S E W Q P E I E E L T N N F Q
54546 agctatcaaaaactcgaaatggatgcaagaggtcgagtgacggttaagcatttccagtcctactcaattagcaattctgtttat
337 S Y Q K L E M D A R G R V T V S I S S P T Q L A I L F Y
54630 gatattcatgggttgaaaagctcctgaaagggttaaaacctagaggaaacaggcgaaagtattgtcgagcattttgataacgatatc
365 D I M G L K S P E R D K P R G T G E S I V E H F D N D I
54714 tcaaaagcacttttgaaatatagaaaatagtttgcacctatacaacacttgaccaacaccttgcaagcctgac
393 S K A L L K Y R K Y A K L V S T Y T T L D Q H L A K P D
54798 aatcgaattcacactacattcaaacagtagcaggttaagacagggcggtatgtcaagtgaagaatcctaactacagaatattcct
421 N R I H T T F K Q Y G A K T G R M S S E N P N L Q N I P
54882 tctcgcggtgaggggtcagtagttcgacaaactttgcagccagtgaaaggcattacattattggtagtgactacttcaacaa
449 S R G E G A V V R Q I F A A S E G H Y I I G S D Y S Q Q
54966 gaacctcggttcattggcggaattaaagtggcgagaaagatgcgacatgcttacgaacaaaacctggacctatattcagttatc
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55302 caggactgggatattgttcaaacagctaccggtcgagaagaagggttctctgataggtcttctgaatacagaggttcgagtat
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40401 atgacaaaaatttatcaactcatacggccctcttctcacttgaacctttacgtcgaaacaagttagtcaggagcytaacgaacaactcc
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85 T V P H N S D G T K T M C S V W A S F D P N N G V H G N I
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225 D S V R P T F S G I S L V D T T S A V R Q I L T G N N F
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 141 R V L M T T V V A N A A C Q I D V Q F Y T S M P C Q F T Y T T V
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367

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 225 C W L T Y A L T D G E S N Q I Y M T E S G Q T T I K E T
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 46976 ctgcttaaatag 46987
 561 L L K *
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421 P L K A K A S R S T A R L R S K V T R E G V E A F *

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1 M T D F K K R F K K A V T E T I N R D G I E N - L M - D - W L
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29 E N D T N F F S S P A S T R Y H G S Y E G G L A G R S A
13328 aacgtgttcaatcaactacttttcgaaatggataccatggttaggcaaaagctgggaagacatttaccatggaaacagttgca
57 N V F N Q L L F E M D T M V G K G W E D I Y P M E T V A
13412 atcgtfagcacttttccagaccttgcgaatgttggtagtcgtaaaactgaaaaatggcgcaagacgacgaggtgaaatgg
85 I V A L F H D L C K V G Q Y R E T E K W R K N S D G E W
13496 gaaagctatttagcatatgaatacgcacctgagcaacttacaatgggacatggtgcaaaatctaatttcttctcaacgcttct

369

113 E S Y L A Y E Y D P E Q L T M G H G A K S N F L L Q R F
13580 attcaactcacgccagttgaagctcaagcaattttctggcatatgggagcctatgatattagtccttatgcaaatttgaatgga
141 I Q L T P V E A Q A I F W H M G A Y D I S P Y A N L N G
13664 tgtggagcagccttcgaaactaatccacttgacttccatccatcgccagatattggccgcaacttatgtatgcgaaatgaa
169 C G A A F E T N P L A F L I H R A D M A A T Y V V E N E
13748 aacttcgaataactctcaaggtccagttgaacaagaggtgaggttgaagaagtagttgaagaaaaacctaagagttcaactcgt
197 N F E Y S Q G P V E Q E A E V E E V V E E K P K S S T R
13832 aagaaacctgcgcctaaaggaagaaagttgaagaggtgaagaaaaaccaaagctggaatcactcgactcgcaaacctgcg
225 K K P A P K E E K V E E A E E K P K A G I T R R R K P A
13916 ccaaaagaggaagaggtagaagagcctaaagaagagcctaaagaagcatcttctaaaattcgaatgcctaaaaagactgaaaag
253 P K E E E V E E P K E E P K K A S S K I R M P K K T E K
14000 gtcgaagaggtagaagcgacagcagcgaaggtgaagaagcagaggacgacaatgtgggtggtacctgtggtatgttcga
281 V E E V E S A D E P K V E E A E D D N V V V P A G Y V R
14084 gatgtctactacttctacagtgagtcgctgacttactacaagaagatgtcgacgagcctgacgatgacgacgacattctt
309 D V Y Y P Y S E V A D V Y Y K K D V D E P D D S D I L
14168 gtacgcaagaagagtacatggacgcaatgtgctcctgtattagaagaagacttctctacgaacttgacggcgaaggttcacaaa
337 V D E E E Y M D A M C P V L E E D F F Y E L D G K V H K
14252 ttacgcaaaaggtgaacgcttgccggaagaatacgcagcaagaagaaacttgggaacctatcactgaagcagaatacatcaagcgaaca
365 L A K G E R L P E E Y D E E T W E P I T E A E Y I K R T
14336 gaaaaacctaaagcagttgcaaacctactcgaaaactccagcgcttctcgtcgccctcgcccttaa 14404
393 E K P K A V A K P T R K T P A P S R R P R P *

dp1ORF010

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29 P R I P F S A P S M N Y Q T Y G G L P R K R V V E F F G
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85 T E E L K E K L E N A R A S K A S K T A V K E L E M Q L
9035 gatagtcctcaagagcctcttaagattgtatatcttgaccttgagaatacatttagacactgagtggtgtaaaaagattggagtc
113 D S L Q E P L K I V Y L D L E N T L D T E W A K K I G V
9119 gatgttgcaaatatttgatagttcgccctgaaatgaacagcgtgaagaaacttcaatattgttttagacatttgcgaaca
141 D V D N I W I V R P E M N S A E E I L Q Y V L D I F E T
9203 ggtgaagttggcctagtagttctagattccttgccttacatggtcagtcacaaacctatttgatgaagagttgactaaaaagggc
169 G E V G L V V L D S L P Y M V S Q N L I D E E L T K K A
9287 tatgcaggaatctcagcgctttagtgaatttagtgaaggttactcctcttctactcgctacaaacttgcattatcttaggc
197 Y A G I S A P L T E F S R K V T P L L T R Y N A I F L G
9371 atcaatcaaattcgagaagatagaatagtcagtaaatgcctattcaactccaggcggaagatgtggaagcagttgtgtgca
225 I N Q I R E D M N S Q Y N A Y S T P G G K M W K H A C A
9455 gttcgacttaaattagaaaaggtgactaccttgacgaaaacggtgcatcattgaccgtagctcgaacacctgcaggggaat
253 V R L K F R K G D Y L D E N G A S L T R T A R N P A G N
9539 gtatagagtcattcgtagaagaccaaagcttaagcgggacagaaaattagtttctctatcagcttctctcattgagga
281 V V E S F V E K T K A F K P D R K L V S Y T L S Y H D G
9623 attcaaatgaaaatgacctgttagatgtcgctgtcgaatttggagtcattcaaaaggcaggggcatgggtcagtagtcgac
309 I Q I E N D L V D V A V E F G V I Q K A G A W S I V D
9707 cttgaaactggagaattatgacagatgaagacgaagaaccattgaagttccaaggcagggcaaatctagttcgacgcttcaag
337 L E T G E I M T D E D E E P L K F Q G K A N L V R R F K
9791 gaggatgactacttattcgacatggtgatgactgcggttcacgaaattatcactcgagaagaaggtcaa 9859
365 E D D Y L F D M V M T A V H E I I T R E E G *

dp1ORF011

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28101 ccaactcttttccctaattgctcaacaaacagggacagacatttcatggctcaaggggtgcaataatttgcagtaactatccag
29 P T L F P N A Q Q T G T D I S W L K G A N N L P V T I Q
28185 ccatctaacgacgcgaaagcaagtcctcgtaacgtgctggatttagcaaaacagctactgagatggcattcttcggtgag
57 P S N Y D A K A S L R E R A A G F S K Q A T E M A F F R E
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85 S M R L G E K D R Q N L Q M L L N Q S S A L A Q P L I T
28353 caactctataatgataactaagaacctttagacggttgtaagcgcaagcagaatacatgcgtagtgaatttgcattacgggt
113 Q L Y N D T K N L V D G V E A Q A E Y M R M Q L L Q Y G
28437 aaattcactgtcaaatcaactaacgagggctcaatacacttacgactacaacatggatgctaagcaacaattgacgtcact
141 K F T V K S T N S E A Q Y T Y D Y N M D A K Q A Y A V T
28521 aagaaatggactaacccagctgaaagtgacctatcgctgacatttttagcagcaatggatgacatcgaaaatcgtaggggtgtt
169 K K W T N P A E S D P I A D I L A A M D D I E N R T G V
28605 cgccctactcgaatggtcttgaaacgaaacacttataaccaaagtgactaagagtgactctatcaagaaagctcttcgaatgggt
197 R P T A R M V L N R N T Y N Q M T K S D S I K K A L A I G
28689 gttcaaggttcttgggaaaacttcttcttgcgaagtcagctgagaaattcatcgctgaaaaaacaggtcttcaaatcgct
225 V Q G S W E N F L L L A S D A E K F I A E K T G I A Q I A
28773 gtctactctaagaaaattgctcagttcgctgacgctgacaaacttctgacgttggtaacattcgctcagttcaacttgattgac
253 V Y S K K I A Q F A D A D K L P D V G N I R Q F N L I D
28857 gacggtaagtggtattgcttccacctgacgcagttggtcacacttggtagcgttactactccagaagcattcgacttgggttca
281 D G A V V L L P P D A V G H T W Y G T T P E A F D L A S
28941 ggcggaacagacgctcaagttcaagttctttagcgcggaacctaccgttacaacttatctgaaaaacatctcgtcaacattgca
309 G G T D A Q V Q V L S G G P T V T T Y L E K H P V N I A

29025 acagttgtatcagctgttatgattccatcattcgaaggaattgactatgtaggagttctcacactaattag 29096
337 T V V S A V M I P S F E G I D Y V G V L T T N *
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1 M S I K F K T E E L S K I V S Q L N K L K P S K L L E I
5430 acaaaactattggcatatttttggtgacggcgatcgctcatgtttacagcgtatgatggctcaaacttcttcgtagcattatc
29 T N Y W H I F G D G E C V M F T A Y D G S N F L R C I I
5514 gacagcgtatgtgaaattgacgtgattgtgaaagcagagcagtttggaactttagaagaaagccacggccgcaacgtgcaca
57 D S D V E I D V I V K A E Q F G K L V E K T T A A T V T
5598 ttagtctcctgaagaatcttcgctaaaagttatgggaatggtgagtagacaatattgatattgttacagaagatgaagagtagccct
85 L V P E E S S L K V I G N G E Y N I D I V T E D E E Y P
5682 acattcgaccacttctcgaagacgtgagtgaaagaaatgctctcactttgaaaagctcgctgtttacggaatcgccaatatc
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5766 aacgattctcggtatcttaaatcaggagcagatggaatttataccggcttctgttaaaaggcggaagcaattactacagac
141 N D S A V S K S G A D G I Y T G F L L K G K A I T T D
5850 atcattcgctatgtatcaacctatcaaggaaaaggactagaatgctcattccttacaacctaatgagtagtttagcaagt
169 I I R V C I N P I K E K G L E M L I P Y N L M S I L A S
5934 attcctgatgagaagatgacttctggcaaatgacgataactgtctatatttcatcggttcggaatctatggaata
197 I P D E K M Y F W Q I D D T T V Y I S S A S V E I Y G K
6018 ttgatggaaggtatggaagattatgaagacgtttcacagcttgactcaattgagtttgaagatgagcggtatccctacagca
225 L M E G M E D Y E D V S Q L D S I E F E D D A A I P T A
6102 gaaatcctgagcgtatttagaccgcttctactattcacttcagcctttgacaaaggaacgctcgaattcttattctgtaaagac
253 E I L S V L D R L V L F T S A F D K G T V E F L F L K D
6186 cgacttcgaatataaaacttctactagcagttatgaagacatcatgtacgcatctgctggcaagaagtttcgaagaagaattc
281 R L R I K T S T S Y E D I M Y A S A G K K V S K E T F
6270 acttgccaccttaacagcttactcttgaaggaaattgtatcaaccgtcaccgaagaaaacttactgtctcttatggaagcgaa
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337 T A I K I S S N G V V Y F L A L Q E P E E *
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1 M N L A S K Y R P Q T F E E V V A Q E Y V K E I L N Q
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29 L Q N G A I K H G Y L F C G G A G T G K T T T A R I F A
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57 K D V N K G L G S P I E I D A A S N N G V E N V R N I I
10467 gaagattctagatacaagctctatggacagcgagttcaagtttacatcattgacgaggttcatatgctttcaaccggagcattt
85 E D S R Y K S M D S E F K V Y I I D E V H M L C G A F
10551 aatcgctgttgaaaaacattagaagagccctcatcggaacccgtgttcattctatgtactactgacctcaaaagattcctgac
113 N A L L K T L E E P S S G T V F I L C T T D P Q K I P D
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141 T I L S R V Q R F D F T R I D N D D I V N Q L Q F I E
10719 agtgaaaaatgaagaaggagctggttatagttatgagcgtgacgccccttctggttattgggaaacttgcaaatggaggaatgcgt
169 S E N E E G A G Y S Y E R D A L S F I G K L A N G G M R
10803 gacagatcacacaaggctcgaaaaagtccttgattatgctcacgcttgacatggaagccggttctaatgcaactagaggttccg
197 D S I T R L E K V L D Y S H H V D M E A V S N A L G V P
10887 gactacgaaacattcgcttcaattgttgaagctattgccaactatgacggctcaaaggtttagaattgttaaatgacttccac
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10971 tactcaggaagaaacttgaattagtgaactcgaaactttacagacttcttttagaggttgaagattggctagttcgagat
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281 I S I T Q L P A H F E S K L E Q F C E A F Q Y P T L L W
11139 atgctagaagaaatgaatgaacttgcgtgaggtgttaaatgggagcctaattgctaaaccgataattgaaccacaaacttctttg
309 M L E E G M N E L A G V V K W E P N A K P I I E T K L L L
11223 atgagcaaggagagtgga 11240
337 M S K E E *
dp1ORF014
50961 atgaaagtaaatggtcttcaaatgaaagcagctcctgaacaaataattgaaaaactttcgagacaacttgaagacgaaggaaca
1 M K V N G L Q I E A T P E Q I I E K L S R Q L E D E G T
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29 F I F R R T K S L G S N Y Q P S C P F H A G G T E K H P
51129 tcttgtggcatgagtagaatacttcttattcaggaagtaaggtagcggaagctggaacggttactgtttcacttgcggctac
57 S C G M S R N P S Y S G S K V T E A G T V H C F T C G Y
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85 T S G L T E F V S N V L G R N D G G F Y G N Q W L K R N
51297 ttggaaacatctagcgaagtagtttaggcaagggctgacccctgaagcgtttcgaaagaaatgggagaactgaaaagtcgagcat
113 F G T S S E V V R Q G V S P E A F R R N G R T E K V E H
51381 aaaaatcattcgaagaggaaacttgataaataccggtttattcattccttatatgtatgaacggaattgacggagaggtcatc
141 K I I P E E E L D K Y R F I H P Y M Y E R K L - T D _ E - 4 I
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169 E M F D V G Y D K L H D C I T F P V R N L K G T V F F
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197 N R R S V R S K F H Q Y G E D D P K T E F L Y G Q Y E L
51633 gtagcatttgcagactattttgaaaaactatttagtaagtagctgagtagctgttatcaactcagtagctgttgggtca
225 V A F R D Y F E K P I S Q V F V T E S V I N C L T L W S
51717 atgaagattccagcagtcgctcttattgggagtaggtggaggaaatcaatttactaaaacgacttcttataagaatatt

371

253 M K I P A V A L M G V G G G N Q I N L L K R L P Y R N I
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281 V L A L D P D N A G Q T A Q E K L Y R Q L K R S K V V R

51885 tttttgaactaccctaaagagttctatgataataagtggtgataaaacgaccatccggaattattaaattttaatgatttagtc
309 F L N Y P K E F Y D N K W D I N D H P E L L N F N D L V
51969 ttgtag 51974
337 L *

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141 E D F K W L N L M L E T T F E G G K H I P Y I G I S P A
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225 G A M G N I M T S K G L V D L S Q K N G G I D A V R R L
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253 P K P V. Q V E I E S I I E E T G A H F S L E Q L V E D Y
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281 K L R A L F N V Q Y M L N W A E N Y E F K G I K N R Q R
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309 R L F *

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57 E Y M H A W L I E N G Y E L I S E N A P W D A K R G D I
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281 K P Q F T V E P D G L I T A K V *

dp1ORF017
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1 M I G Q G L V K S T I S K W K Q L P K Y I I V E G E V G
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141 V D T S G I D D R A I V D Y C N L A S N L Q M L E D L
11746gaatatggcgagaaagactatttgaagggttacaacattttatgacttaatatgggaggcaagtgctaggaatttcgctaaag
169 E Y G A E E L F E K V T T F Y D L I W E A S A S N S L K
11830gttactaattggctcaaatttaaggaaactgatgaaggaataatgagcctaaacttttctcaactgcttttaaatggctcg
197 V T N W L K F K E T D E G K I E P K L F L N C L L N W S
11914acagttgtcatcaggaagcactatgaaatgcttctgaagaacttgaggccatgaccttttagtgagggaagcatcagg
225 T V V I R K H Y V E M S F E E L E A H D L L V R E A S R

372

11998 tgtttgcgaaaggatctctaaaaagggtcaaatgcgcgtgtctgcgtgaacgaatttatcaggagggtcaacaagttgagtga
12081
253 C L R K V S K K G S N A R V C V N E F I R R V K Q V E *
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35847 atggctagcagacagcgtatttggtcgacggaattgacctgtcgacaaagggtgcaaccgtgctagaatatgtaggactcact
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35931 ttccgaggtttaaggactcaggatttaaaacccctgaaggcatagacggagtattagattctccgtctaagctatgtccgct
29 F A G F K D S G F K N P E G I D G V L D S P S N A M S A
36015 cttactggaagcgtgaccttaattgttccacggagaaaccgaaagcaagtttaacaaaaacagggcagttcaacaatttatt
57 L T G S V T L M F H G E T E K Q V N Q K Y R Q F K Q F I
36099 cgctcgaagtcattttggagaatttcgacacttgaagaccctggatactatcgaaacgggaaaatttttaggagaaaccgagcaa
85 R S K S F W R I S T L E D P G Y Y R T G K F L G E T E Q
36183 ggaaaacttgtagcgttcaagcctttaagatacttcccttgtagttaattagggattcagttcaaatgcttaccaggtac
113 G K L V D V Q A F K D T S L V V K L G I Q F K D A Y E Y
36267 agcgactcaactgttcgaaagggttataagttcaaccgcctttggaggcgatagcttaccacccaggaagacctactcga
141 S D S T V R K V Y K F Q P A L G G D S L P N P G R P T R
36351 caatttagagtgaataaagaactacttctcaaatcaaggatattttcgaattggcgaaaaaagttcaggacagttgtttgag
169 Q R Y G T L R L M V T K I D K R S K L L K A F P D N C V
36435 ttccggtactaattcagttatgtatggaagtggtcgattattattctaaatcttggaacttttgaaactattaaaattagcagt
197 F G T N S V L M E S G S I I I L N L G T F E L I K I S S
36519 gcaaatcaagcgactaacttatttagatacattaaacgagggcagcttctcaagattcctaagtaaattcaacaattaccatt
225 A N Q A T N L F R Y I K R G A F F K I P N G N S T I T I
36603 gaataccgagccgatgacgcagcagcttggacctctactctcccgctcaagttgaactgtttctaaatccgtcttactattag
36686
253 E Y R A D D A A A W T S T L P A Q V E L F L N P S Y Y *
dp1ORF019
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1 M N V Y L N Q M G N V V R E T S V S T V W K T L T Q K G
12245 ctccgtttctaatcatcgatattctcgtgttcgagatgataaggagtttctgtctaatgagtcgaggtggaaaaggcttcggat
29 L V S N H R I F A V R D D K E F L S N E S R W K R L P D
12329 gttagatatgggacacttgttttgatggttactaaaaatgacaagcgaagcaagttgctaaaggcctttctctgataattgtgtt
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85 E F E K M T D A Q L K R H F V S K Y S T I D S D M I D M
12497 gttatccagttctgtctaaacgactctagaattgacaatgaattggacaagctgtcgagattgaaaagggttgacgcatca
113 V I Q F C L N D Y S R I D N E L D K L S R L K K V D A S
12581 gtggttaatccattgtcgaagcacaagaccgaattgacattttcagcctagttgatgattggaataataggccgagcag
141 V V E S I V K H K T E I D I F S L V D D V L E Y R P D M
12665 gcaattatgaaagtactgaacttttagccaaaggagaaagtcctattggattgcttaccttgctttatcaaaattttaataac
169 A I M K V T E L L A K G E S P I G L L T L L Y Q N F N N
12749 gcttgcctgtgctaggagccgatgagcctaaagaagccaatttaggcattaaagcagttcttaatacaatattgtctataac
197 A C L F N D K G E Q I C N H V T L T G G N P A L N E P
12833 ttcaatacagagctggactcagccttgaaggcatggctatttttaggtcaagctatcgagggcataaagaatgggtcgctataca
225 F Q Y E L D S A F E G M A I L G Q A I E G I K N G R Y T
12917 gaaagttcagtggtctatatttcttggatataaaatttttctacttacttaa 12967
253 E S S V V Y I S L Y K I F S L T *
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29 F G P T I Q G E G M V I G Q K T I F I R T G G C D Y H C
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57 N W C D S A F T W N G T T E P E Y I T G K E A A S R I L
2116 aaactagctttcaatgataaagggtgaacagatttgtaaccacgtgacattgactggaggaatcctgccttaatacaacgagcct
85 K L A F N D K G E Q I C N H V T L T G G N P A L N E P
2200 atggctaaagatgatttcgattctaaagaacattggattcaagtttggtctcgaaactcaaggaaactcgattccaagaatgggttc
113 M A K M I S I L K E H G F K F G L E T Q G T R F Q E W F
2284 aaagaagtgaagcgtatcactattagtcctaaacgcttcaagtggaatgagaactaatatgaaaattcttgaaagctattgta
141 K E V S D I T I S P K P P S S G M R T N M K I L E A I V
2368 gatagaatgatgaaaaccttgactggcttttaaaatcggttatctttgacgaaaatgacctagcttatgagcggtgatag
169 D R M N D E N L D W S F K I V I F D E N D L A Y A R D M
2452 tttaaaactttcgaaggcaagttacgtccagtgaaactacctttcagttgggaatgcaaacgcatacgaagaaggaaaaatcagt
197 F K T F E G K L R P V N Y L S V G N A N A Y E E G K I S
2536 gataggtctcttgaaaagttgggtgggtttgggataaagtgatgaagaccagctttcaacaatgttcgacatttaccgcaa
225 D R L L E K L G W L W D K V Y E D P A F N N V R P L P Q
2620 ctccatcacacttgtttatgataataaaaggaggtataa 2658
253 L H T L V Y D N K R G V *
dp1ORF021
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29 Q L S T M F D L Y R N F I H L F M I I K E E Y K M K I E
2672 catctagataaaatcggttaacgtatttagggagagagaacggatgggttcccttaagccggatgaaattgttaaccttggaacaat
57 H L D K I G N V L G R E N G W A S L K P D E I V T A R D M
2756 actgaggcagccgttcaaaagacttttgggtctattaggcgaggacgcagaacgtgacgggttgcaagatactccattccgtttt
85 T E A A V Q R L F G L L G E D A E R D G L Q D T P F R F

373

2840 gttaaagcactcgctgaacataccgtagggtatcgagaagaccctaaacttcatctcgaaaaacattcgagctcgaccatgaa
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2924 gaccttgttctgtgaaagacattccattcaattctttatgtgagcatttagctccgttcgtagggaaggtgcatattgca
141 D L V L V K D I P F N S L C E H H L A P F V G K V H I A
3008 tacatttctaaggataagattacaggtctttcaaaattcggtcgagtggttgaggatacgttaaaccgacttcaagtacaagag
169 Y I P K D K I T G L S K F G R V V E G Y A K R L Q V Q E
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197 R L T Q Q I A D A I Q E V L N P Q A V A V I V E A E H T
3176 tgcattgagcggacggttattaagaagcacggggcaacgacagtgacttcaactatgagaggtcttttccaagatgacgcattc
225 C M S G R G I K K H G A T T V T S T M R G L F Q D D A S
3260 gctcgagcagaattgcttcagttgattaaaaagtag 3295
253 A R A E L L Q L I K K *

dp1ORF022

30896 atgagtaagacattctttacggaatcaagctcggtgcaaatcgaggagcttgaccattgactcagttgcaaaagtcggcgga
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30980 gctaacttgtcgtagatacggcagaaacagcagaactcgaagccgtgacctcggagggaactgaagatgtgaaacgcaatgac
29 A N F V V D T A E T A E L E A V T S E G T E D V K R N A
31064 acgcgcattcttgcattcgctgactccagacctttatacgggttatgacttaacattcaaggacaacacggttgacctgaa
57 T R I L A I V R T P D L L Y G Y D L T F K D N T F D P E
31148 atcatggccctaattgaagtggtacagtcagtcacaaagcggaactattgctggatacgcacaccccaatgcttgacaaaggt
85 I M A L I E G G T V R Q Q G G T I A G Y D T P M L A Q G
31232 gcttctaataatgaaaccatttagaataacacattatgtgccaactatgtaggtgactcaattgtcaactacgtgaaaatcact
113 A S N M K P F R M N I Y V P N Y V G D S I V N Y G K I T
31316 ttgaataactgtaccggttaagctccagggcttcaatcgggaaagagttctacgctcctgagttcaacatcaaggcacgtgaa
141 L N N C T G K A P G L S I G K E F Y A P E F N I K A R E
31400 gcaaccaaagcaggtttgccaagttaagtcaggtatgtggcacaacttccagcggttcttcgtcgcggtgacattcgattg
169 A T K A G L P V K S M D Y V A Q L P A V L R R V T F D L
31484 aacggtggaacaggaacccgacgcagtcgagttgaagcaggtgaagaagatttctccaaacagtgacctacaccttaaca
197 N G G T G T A D A V R V E A G K K I S P K P V D P T L T
31568 ggtaaggtcttcaaaaggtggaagttgaaggtcaactatttgggacttcgacaaccacatgctgacctgacgcagctc
225 G K A F K G W K V E G E S T I W D F D N H M M P D R D V
31652 aaactcgtagcacaatttgcatag 31675
253 K L V A Q F A *

dp1ORF023

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6503 ctgaccgggttattgaacgaactcagcctgaatataatccttcgacataattataagccagcgggttggtggatgtattcga
29 L T R V I E R T Q P E Y N P S T Y Y K P S G V G G C I R
6587 aaaatgtatttgaagaatcggtgagtcattatagataacgcagatttcaacctaattgcaatggcggaagctggaacattt
57 K M Y F E R I G E S I I D N A D S N L I A M G E A G T F
6671 aggcacgaagttctccaaggtacatggttaaaatggctgaatcgatgaggacttgaaatggttgaaatgtagcagagttcttg
85 R H E V L Q E Y M V K M A E I D E D F E W L N V A E F L
6755 aaagaaaaatccagttgaaggaactatcgctcagcagcgttcaagaaaaagattgaacgaagtgtaagaacgaacttctt
113 K E N P V E G T I V D E R F K K N D Y E T K C K N E L L
6839 caactttcattcttgtgtgacggactagttcgatataaaggcaagctctacatttttagagattaagactgaacacattgttcaag
141 Q L S F L C D G L V R Y K G K L Y I L E I K T E T M F K
6923 ttactaaacatactgagccctatgaagaacacagatgcaagcaacttgctacggaatggtctagagtcgatgatgctatt
169 F T K H T E P Y E E H K M Q A T C Y G M C L G V D D V I
7007 ttcctttatgaaaaatcgagataactcgaaaaagaaagcctacacgtttcacatcacagacgagatgaaaaatcaagtccttgga
197 F L Y E N R D N F E K K A Y T F H I T D E M K N Q V L G
7091 aaaattatgacctgcgaagatgtatgagaaaagcgaaagtcctaaatctattgctcttcagcctattgccatattgtaga
225 K I M T C E E Y V E K G E S P K I Y C S S A Y C P Y G A C R
7175 aaggaaggtcgaaatctgtga 7195
253 K E G R N L *

dp1ORF024

25992 atgaacgcagtagatggccaggtgattcatattctacaagtattagcagaagatggaaatgctacggtgaaaagttcgaaaag
1 M N A V D G Q V V H I L Q V L A E D G N A T A E K F E K
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29 E V R A A S L V F S R R A A E A V V K G E I Y K D G K N
26160 ctctcgaacgtgtttggtcttcagccgcagcgaggaatgatgttcaacaaatagtcacacaaggcctagcaagtggaatg
57 L S K R V W S S A A R A G N D V Q Q I V T Q G L A S G M
26244 tctgctacagatatggttaaaatgctcgagataatatcgaccctaaggttcgaaaagattgggactttgataagatgctgag
85 S A T D M A K M L E K Y I D P K V R K D W D F D K I A E
26328 aagctagggaacacgtgctgctcataaatatcaaaatctgaatacaatgcccttcgacttgctcgaactaccattagccattcc
113 K L G K P A A H K Y Q N L E Y N A L R L A R T T I S H S
26412 gccacagctggagtgagacaatggggcaaggttaacgttattcgctcgaaaagttcaatggcattctgctcagcgtccaggtcga
141 A T A G V R Q W G K V N P Y A R K V Q W H S V H A P G R
26496 acgtgtcaagcgtgtatcgatttagatggtgaagtatttctatcgagaatgtcctttcgaccatcctaatggaatgtgctac
169 T C Q A C I D L D G E V F P I E E C P F D H P N G M C Y
26580 caaactgtatggtacgaaaactcactcgaagaatcgctgatgagttgagaggtgggtgagcggagaacctaagtgattatta
197 Q T V W Y E N S L E E I A D E L R G W V D G E P W D V L
26664 gacgaatggtacgacgatttaagttcaggaaggttgagaaatacagcgacctcgacttgtttaaagttattag 26738
225 D E W Y D D L S S G K A G E K Y S D L D F V K S Y *

dp1ORF025

18778 atggcaagaacaaaaagcgaaaaaagtaaatgtcaaaaggaaaatgcttatccctacaaatctctcgaaaaaagtaaatgta
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18694 aaagcaatcgcttatagaaaagtcactgttaagtggctgcctaaacacagatgaaattcaagtatatatttcgacctttatataaat
29 K A I A Y R K V T V K W L P N T D E I Q V Y F D L Y I N
18610 aaaaacaggctgacaatgtaggcactattgacccggacaagagctatgttgaaggaattaggattgttgaagaacactcag
57 K N R L T M L G T I D P D K S Y F E G I R I V C K K P Q
18526 ccttgatgactgttaaggagctccagggtgcgcgtgcagacgccccagggttttttgcagttcttaagcctattgtcacacg
85 P W M T V K E L Q V A R A D A P G F F A V L K A Y C H T
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113 V G D V L D S G A E P T E I V Q G I M Y K D G E L F K D
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141 S E I V S L F K Y D V K E P Y E F P K D L P I T L D N F
18274 ttagagttcattatgtctagccagcactagagcacttgttttgcgttgcgttaatataggtgagttttccaagaattggcgg
169 L E F I M S S Q H T R A L V L R C A N I G E F S K N W R
18190 aaatggcaaaaagctatccagactcctgcgtactatgccaaggcggtgactttaaagtagacgaaactgtttgggacttttca
197 K W Q K A I Q L L L D Y A K A D D F K V D E T V W D F S
18106 cccggctctaaagctggaaggtagcagctcgtaagggtatgaggcaattcaacaagccttgagcagataataataa
18026
225 P G S K A G K V A R R K G Y E A I Q Q A L E Q I N K *
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21512 atggcgaaagctactggacaaaagttcgaagaggaaaaactcctccagcgccaaaagacaaaaaggaatcaaagcaaatgcg
1 M A K A T G P K V R R G K T P P R P K D K K G I K A N A
21596 cgtgtcaataaagaccagttcgtagagtactataaaggcatcaagatgacaattaaaggaacgtgactagaatgaaattg
29 R V N K D Q F V E Y D Y K G I K M T I K E R D A R M K L
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57 E F I R G M T I Q E I A A R Y G L N E K R V G E I R A R
21764 gataaattgggtgaaggctaagaaaaggttcgagaatgaaaaggctcctgttactaatgatacattgactcaaatgtatgcaggg
85 K W V K A K K E F E N E K A L V T N D T L Y A G
21848 tttaaagctcagtcataatataatcacgcgcctgggagaaactaatgaacatcgctcgaatgtgtttagataatcctgac
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141 R Y L F T K E G N I R W G A L D V L S N L I D R A Q K G
22016 caagaagagcggaatggaatgctccggaagaggttcgatataagatacaaatgagcgcgagaaaactacattgtctccggg
169 Q E R A N G M L P E E V R Y R L Q I E R E K I T L R A
22100 aaaatgggagaccaggaattgaaggcgaggttaagataacttcgtagaagcactagataaagcagctcaagccgtttggcaa
197 K M G D Q E I E G E V K D N F V E A L D K A A Q A V W Q
22184 gaatttagtgacgcaacaggttctacattaaggagtgactgataatgacaataagcctgagaaataa 22252
225 E F S D A T G S Y I K G V T D N D N K P E K *
dp1ORF027
52762 atgggaaaagttatcaattcaaaaacaggaaacttttagctcagggtctaataacagagtttttcacactcgctgaccacggtgac
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29 S A I V T L L Y D D P E G E D M D Y F V V H E A D V D G
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57 R R R Y I N C N A I G E D G E T V H P D N C P L C Q N G
53014 ttccctcgattgaaaaactatttctcaactttacaacatgatacgggaaaaaggtgaaacatgggacgagggcgtttctat
85 F P R I E K L F L Q L Y N H D T G K V E T W D R G R S Y
53098 gttcaaaagattgttacatttatcaataaatatggaagccttgtagtactcagccttttgaaattattcggttcaggagctaaagg
113 V Q K I V T F I N K Y G S L V T Q P F E I R S G A K G
53182 gaccaacgaactcttatgaattccttcagagagctcgggaagacagtgactactcctgaagattttccagaaaagagcgaactt
141 D Q R T T Y E F L P E R P E D S A T L E D F P E K S E L
53266 ctgggaactctaatttagacctcgacgaagacaaaatgtttgacgtggttgacggcaagttcactcttcaagaagagcgttct
169 L G T L I L D L D E D Q M F D V V D G K F T L Q E E R S
53350 tcaagtcggttcaaatcacgtagaggagcatctcctgcgcctagacgaggttcgggtcgagaatctcacaaggtcgaaacagct
197 S S R S N S R R G A S P A P R R G S G R E S S Q G R T A
53434 gaaagaactccttcagttcgaaagaactcctccaacacgaggtcgaggattctaa 53490
225 E R T P S V S R R T P P T R G R G F *
dp1ORF028
44595 atgtcaaaaattaaattcgaaaaccttaaaaaaggcgatgttgtgctacgagctaaatctcaacgaagtttaaatcggttca
1 M S K I K F E N L K K G D V V L R A K S Q T K F K I V S
44679 atttttagcagacgaaaagaaagcagaccttgaatcattagaagacggaggtgaacttcacctttcagcttcaactctcgaacgt
29 I L A D E K K A D L E S L E D G G E L H L S A S T K A L
44763 tggtagacaaatggaagatgaaactgaaccttaaaaaagaagaagctgctaaccctgcttaaaaaggctgctcctcagttgtcga
57 W Y T M E D E T E P K K E E A A K P A K K A A P A V A R
44847 cctgctcgaaaaggtagagtcggtcccaaaccttaaaaaagaagtccttgaggaagaatctcgaagtttaaggaacagcgga
85 P A R K G R V V P K P K K E E V L E E E I P E E Q P E
44931 gaagttgggtcagttagtgagaaatctactgttcgaaaacctgctccttaaaaaagaagcgtgatggcgattactaaggctctt
113 E V G S V S E K S T V R K P A P K K E S V M A I T K A L
45015 gaaagtcgaattgttgaagcctttcctgcgtcactcgaactcgtcactcctacatcgctccttaagaagaacttc
141 E S R I V E A F P A S T R I V T Q S Y I A Y R S K K N F
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45183 tctattgtcctcgtcattacgaatggcgagtagcggaatttttaactcgtaaggagaagatttgacacccgaatggaa
197 S I A P A S Y E W A I D G I F K L V K E E D I D T A M E
45267 ttgattgaagcttctcacctttcttcgctatga 45299
225 L I E A S H L S S L *
dp1ORF029

20088 atgaataacgaaaaaattattgaaaaaattaaaaatcttatttcaattagcaaatgacaaccggagtgacgaagaggggcaaat
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 29 A L L M A Q K L M L K N N I A L A Q V E Q F D E P K Q F
 19920 gagactctcaagctgttgggaagaagcaggtcgaaatattttgggtgggaacgtgaacttggtcatattctcgcgactaat
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 19836 aggtgcttttgattaatcagcgtgatatgcgcttgaataaaaagtcgaataattttcttcgcgcaaaaacaagacgctgaatta
 85 R C F C I N Q R D M R L N K K S R I I F F G E K Q D A E L
 19752 gtgtctaaaaatgatgaggtcgttctgttattcttcgttaccgtatttgaccgacttctactcgcgaaccttctcacaagaat
 113 V S K I Y E A A L L Y L R Y R I D R L P T R E P S Y K N
 19668 tcatactcaaaggctttttgtcagccttgacctgatttataaaagcaggtggaagaatttcaatttggtctcacttagc
 141 S Y L K G F L S A L A I R F K K Q V E E Y S L M V L P S
 19584 gagcaaacaaaaatgcgcttcaggacacatttcgaaatttaaagaaggaaggaattgacagacctcaacgatcctcaatctt
 169 E Q T K N A A L L Q D T G F R N L K K E G I D R P Q H D F N L
 19500 gaagcgtatattgaaggcggtttcctggcgagaatgcaagaattgcccgatgaattttggaaggggtaactaa 19423
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dp10RF031

26943	atggcgttatcaattagaagacttggtaaaaggctctagatgaaccaactatcaaacagggtgaaggaaattatttcgaaaacttcg
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29	aaagaactcgatgctaaaaatttcttgatgcgcgcagctcaacacttgacctcacgcacgttctgatgaagttgttcaacag
27111	K E L D A K I F I D G D G G Q H F V P H A R F D E V V Q Q
57	cgcgatcgactacgggctcaattaattcttataaagaacaagtcgcgcagcgttcttaaacagggtcaaagataacgggtgatgcg
27195	R D A A N G S I N S Y K E Q V A T L S K Q V G K D N G D A
85	cagacactatccaaaaccttcaagagcaactcgacaagcagctcgaactgcgaaggcgctgtgattacttcagctcttcat
27279	Q T T I Q N L Q E Q L D K Q S Q L A K G A V I T S A L H
113	ccgttgattagtactccattgctccagcagcagacacttctggattatgaaccttgacaactacgggtcgaaagtgacgggt
27363	P L I I S D S I A P A A D I L G F M N L D N I T V E S D G
141	aaagctaaaggctcttgatgaagagttgaaaggctgttcgtgagtcctcgtaataacttattcaagaagtcgaagttcccgagaa
27447	K V K G L D E E L K A V R E S R K Y L F K E V E V P A E
169	caagaggctcaagctaaagtcggcgcgggactggaatttaggaattccagggtcgtgtcgggtgggtgttcccgacacctcgt
27531	Q E A Q A K A K S P A G T G N L G N P G R V G G G V P E P R
27611	gaaatcggctcttttggtaagcaactgtgctgctgctcaacaaacggcaggagcacaagaacaatcattctttaaataa
197	E I G S F G K Q L A A A Q Q T A G A Q E Q S S F F K *

197
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52201 cgacatgacattgaaagcattgccttcagactattcctcaactgtttggcaacgcttcaaatcaaaccaaggggccaagcttttca
57 R H D I E S I A F E T I S K C L L A T F K S N Q G A K F S
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52369 gtgaagtgcagcttcgatagcgttttcgacaaatgaagaaggcgacgattttagtagtctcctatgcacagttggctattgtgaagac
113 V E V T F D S V S T N E E G D D F S I L S T V G Y C E D
52453 tacggaaaaattgaaattgaagcaagcttgcattctatgcagcttttctaatacagagtatgctttatatctcgctctcattcaa
141 Y G G K I E I E A S L D F M T L S N T E Y A Y I S S V I Q
52537 aacggtccttcagtaagcgacgcagaaattgcgcgtgaaattggagtaagcagggtctgctattagtcagtcctagaagtcacta
169 N G P S V S D A E I A R E I G V S R S A I S Q S K-K-S L
52621 aaaaaataattaaagattttatataa 52647 -
197 K N K L K D F I *

137
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1 M A R P K L P Q I D I R E E E I R D A Q D V A D S Y G A
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376

29 I I N K V V D E I V E A A C G S L D Q A M E E I Q I V V
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57 S Q N P V I M E D L N Y Y I G Y L P T L L Y F A A D R A
7922 gaaatgggtggaatacaaatggattcaagttctgtctatcaggaaagaaaaatcagataatctatacattttagccgccgggaaa
85 E M V G I Q M D S S S A I R K E K Y D N L Y I L A A G K
8006 actattcctgacaagcaagcagaactcgaaaacttgctcatgaatgaagaagtcacgaaaatgcttacaagcgagcctacaag
113 T I P D K Q A E T R K L V M N E E V I E N A Y K R A Y K
8090 aaagtccaattaaagctagaacaggccgataaggtattagcatctttaaaccgaattcaaacctggcaactagcagagttagaa
141 K V Q L K L E Q A D K V L A S L K R I Q T W Q L A E L E
8174 actcagtcacaataattcaaaaggagtattattaaatgcaaaaagacgtagacgtgaaaatgattga 8239
169 T Q S N N S K G V L L N A K R R R R E N D *
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1 M S Q N T T R T D A E L T G V T L L G N Q D T K Y D Y D
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29 Y N P D V L E T F P N K H P E N N Y L V T F D G Y E F T
299 tccctttgccttaaaacaggacagcctgacttcgcgaatgttttcattagttacattccaaacgaaaagatgggtgaattctaaa
57 S L C P K T G Q P D F A N V F I S Y I P N E K M V E S K
383 tcattgaaattgtacttactcagtttcctgaacgggtgacttccacgaagattgcatgaacattattttgaattgacttgat
85 S L K L Y L F S F R N H G D F H E D C M N I I L N D L Y
467 gaattgatggaacctaagtacattgaagtcagggcctattcactcctcgtgggtgaatttcaatttaccattcgtcaacaaa
113 E L M E P K Y I E V M G L F T P R G G I S I Y P F V N K
551 tgaattcctcaatttgaactcctgaacttgaacagcttcaacttcaacgaaaattgaacttcttggaatgttcaaggctctt
141 V N P Q F A T P E L E Q L Q L Q R K L N F L G N V Q G L
635 ggacgagctattcgatag 652
169 G R A I R *
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17425 atgcacctaataagagattcgaagatgttgaggacatggaagtccttagcattcaggttcgaaacgaaggtgaggacgacgagt
1 M H L M K D S K M L R T W K S L A F E F E T K V R T T S
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17257 catactgaagctgatattgactacaaagatattctaaattttgtagcttctcgaactcctcaactcgaattcgaactcact
57 H T E A D I D Y K D I L N F V A Y R N S P N P Q I Q I T
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85 S W N A L L S C Y T R N E L S Y K G V S I T D F F E A I
17089 caaactattgcaagttccttcactcacctagactcgaaaacaattgatacacaataatgaaaagcgttcaaacctcgaaggtgaggaa
113 Q T I A S S F T H L D S K T I D T Q N E K R L E R I E E
17005 cttcagtcagaataggtcattgtaactgtactatcgacgaacttaaaaaaggagtcacgaatgcccgatattgaatcagct
141 L Q S R I G H C N C T I D E L K K G V H E M P D I E S A
16921 atttcttaccagtcaggacagattcttgcttgaagatgaacttaattttctgctaaactaa 16859
169 I S Y Q Y G Q I L A Y E D E L N F L L N *
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29 G L S I V I A S N T V G N G K T S W A V R L L Q R Y L A
48976 gaaactgcacttgacggaagaattgttgagaaggaatgtttgtagtgtcagctcaactattgactgagttcgcgactataat
57 E T A L D G R I V E K G M F V V S A Q L L T E F G D Y N
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85 Y F Q T M Q E F L E R F E R L K T C E L L V I D E I G G
49144 ggttccttaaccaaggcctcttattccttctgtatgacttggttaattatagggttgacaataacttgcactatttatacg
113 G S L T K A S Y P Y L Y D L V N Y R V D N N L S N
49228 actaattatactgacgatgaaattattgaccttttaggccaaaggctttatagtcgtatataatgatacttcagtggttctagat
141 T N Y T D D E I I D L L G Q R L Y S R I Y D T S V V L D
49312 tttcaggcaagcaatgtaagaggattggaggtaagcgaattgaatcatag 49362
169 F Q A S N V R G L E V S E I E S *
dp1ORF037
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29 E P L N V K G I L I P P S S W F M G F T F L L I N L I S
56023 aagtcagagaagccaaaatttgcaggttctttgatattgggttaggttattccttacctcgttattgctttatgcaaaaacctta
57 K Y E K P K F A G S L I W V G L F L T S L I C F M Q N L
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85 P Q S L V V A S G V A F W I S Q K A S V F I F D K L S N
56191 aaattagactcgaagattgcaaatgctttgtctagcaacatcggttctattatagacgcaacatattgatttcttaggactg
113 K L D S K I A N A L S S N I G S I I D A T I W I S L G L
56275 agtcctcttggaattggaacgggttcataatagatattccgtcagccgtactaggccaagttcagttcagtttattcttcagcag
141 S P L G I G T V A Y I D I P S A V L G Q V L V Q F I L Q
56359 tcaattgcttcgagatatttgaaaaagtag 56388
169 S I A S R Y L K K *
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1350 atgagagtttctaaaaccttaacattcgacgcagctcatcaactagttggacattttggaaaatgcccgaatttgcacgggcat
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29 T Y K V E I S L A G G T Y D H G S S Q G M V V D F Y H V

377

1518 aagaaaaatcgaggtacattcattgacagacttgaccacgctgttcttcttcaagggatgaacaaatcgcttttagcaaatgca
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1602 gttgacaccaagcgagttctattggatttagaactacggtgagaaatgtcaagattccttacctggacttccacggagctt
85 V D T K R V L F G F R T T A E N M S R P L T W T L T E L
1686 atgtggaagcatgctcgatcgactctatcaaaacttgaggaaactcctacaggttgcgagaatgtactactacgagattttc
113 M W K H A R I D S I K L W E T P T G C A E C T Y Y E I F
1770 acagaagacgagattgaaatgttcaagaacgtaacctttatcgacaagacgaaaagattactgtccgcgaaatttttagagcag
141 T E D E I E M F K N V T F I D K D E K I T V R E I L E Q
1854 gagcaggataatggttaa 1871
169 E Q D N G *

dp1ORF039
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1 M N K S A T F W L V R T A L I A A L Y V T L T V A F S A
3390 attagttatggacctattcaatttagagtcagtgaaagccttgattcttctacctttatggaacctagatggactccggggatt
29 I S Y G P I Q F R V S E A L I L P L W N H R W T P G I
3474 gtattaggaacaattattgcaaaacttctttcactcttggaactgtgacgttttattcggttcactgtactcttctgga
57 V L G T I I A N F F S P L G L I D V L F G S L A T F L G
3558 gtatgggcaatggtgaaagtgtcaagatggcaagtcctctatattcacttatctgtccagttcttgcctaatgcttacttatt
85 V V A M V K V A K M A S P L Y S L I C P V L A N A Y L I

3642 gcgctggaacttcgaatagtttactctttaccttttgggaatctgtcatctatgttaggaattagtgaaagcattatcggttta
113 A L E L R I V Y S L P F W E S V I Y V G I S E A I I V L
3726 atttcatacttcttatttccacgctggcgagaacaatcattttagaacactgataggagcgaaaaatgggatttaa 3803
141 I S Y F L I S T L A K N N H F R T L I G A K N G I *

dp1ORF040
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1 V S Y T G K M F E E D F F E G A K D F E K D A F T V R L
7276 tatgataccactaatggatttcgaggagttgcaaatccctgcgatttatagccgcaactaatttgggaccttgtttattgaa
29 Y D T T N G F R G V A N P C D Y I A A T N F G T L F I E
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57 L K T T K E A S L S F N N I T D N Q W F Q L S R A D G C
7444 aaatttattctcgccggaatttttagtgatttccaaaagcatgaaaagatttatatgggtatccaatttcaagccttgaaaaaat
85 K F I L A G I L V Y F Q K H E K I I W Y P I S S L E K I
7528 aaacggtctggagttaaaagcgtcaacccaaacttcatcgatgcagggtatgaagtttcttacaagaagcgtcgaaactagattg
113 K R S G V K S V N P N F I D A G Y E V S Y K K R R T R L
7612 accattccttccaaaatggttcagatgcagttgagcttattacaaggaagcaaatggcaagacctaa 7683
141 T I P F Q N V L D A V E L H Y K E K S N G K T *

dp1ORF041
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1 M Q K D V D V K M I D P K L D R L K Y T G D W V D V R I
8292 agttctatcactaaaattgacgcccagacgcccagatgtctcaagatgtcgaaaagtgttcaaaaggctcaagtatattcagtg
29 S S I T K I D A D S A D V S R C R K V L Q K A Q V Y S V
8376 gcggcaggtgaatgcattaaaattgcacacggttcttgccttgaaacttctcaagggataggaacatttgcacctcctggttc
57 A A G E C I K I A H G F A L E L P K G Y E A I L H P R S
8460 agtctttttaaagaaaactggtcctaattcttgcctttagcggagtgattgacgaaggttcaaaaggtgacactgatgaatggttc
85 S L F K K T G L I F V S S G V I D E G Y K G D T G E W F
8544 tcagtttggtatgctactcgtgacgcagatattcttctacgaccaaagaattgccaatttagaattcaggaagcaacactgct
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8628 atcaagttcaatttcgtagaattctttaggaatcggttcggtgagccatggaagtagcaggtgatttctaa 8699
141 I K F N F V E S L G N A A R G G H G S T G D F *

dp1ORF042
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57 Q F E P N M K Q V S S F F I V Q Y E F I F N I K C I D Y
48334 aactggttcaacttttcgagcactatgaaaaatgttcgaacttatttaaacattgagtcgaacattgaactttgtcgattttta
85 N W F N F S S T M K N V R T Y L N I E S N I E L C R F L
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113 A E S F V K Y E N V R K R L N L S E R F I T V S T F K R
48502 gcctggatttttgacgaactcgaaggaaaacgggttcaaaattcgaaggattttattag 48561
141 A W I L D E L E G K T G S K F E G F Y *

dp1ORF043
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29 P I H V K I R A A G V M N L I A N G K I P N T L L G K V
31867 acagactgttttggaagaaacttcgacagtcactaaagacaatgctagtcttagcatcaattactgaccaacagaagaaagagcg
57 T E L F G E T S T V T K D N A S L A S I T D Q Q K K E A
31951 ctgcagcgattgaacaaaaccgataccggtattcaagacatggctgaacttcttcgagttatcgagaagettcaatggtagag
85 L D R L N K T D T G I Q D M A E L L R V F A E A S M V E
32035 cctacttcagctgaagtcggcgagtatatgacagatgagcaacttatgacaactcttcagtgcaatgtacgggtgaagtgactcaa
113 P T Y A E V G E Y M T D E Q L M T I F S A M Y G E V T Q
32119 gctgaaacctttctacagacgaaggaaatgtctaa 32154
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378

dp1ORF044

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25582 gttttcaatttcttctgtttctacataaagtgtggaaccgataatggcacttaggacatttcgaagaatctccactctacgccctt
29 V F N F L V S Y I S G E P I M A L R T F E E S P L Y A L
25498 ttcgatatgttttcgaataatctgttttagatgtaaggtcgaacttatgctcacaatggtcacaatcaacttgaacgtctgggt
57 F D M F R N N L F R C K V E L M L T M V T I N L E R L G
25414 cgactccttcttctgggtgtgttcagttgttttcttctgtcatcaacttcgttcttctcactcgtttcatcttgaggct
85 R L L L R L V V Q F V L F L C H Q L R L L H S F H L E A
25330 cctctgttctgtttaaattcgtttgttaatacaggcaatgtccagctgagatttcgtcaagctgagcaagttcttccaaaatgc
113 P L V R L I R L L I Q A M L Q L R F R Q A E Q V L P K C
25246 gttcccatccttctgcgccttttcttcttactga 25211
141 V P I P C P P F P S Y *

dp1ORF045

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1 M K R V K K T K L M T K K N K L N N Q P K K E S T Q T
25424 ttcaaggtaattgtgaccattgtgagcataaagtcgaccttacatctaacaagattatttcgaacatctcgaaaaaggcgta
29 F K V N C D H C E H K F D L T S K Q I I S K H I E K G V
25508 gagtggagattcttcgaatgtcctaagtgccattcgggtcaccacttatgttaggaacaaggaattgaaaaccttattcga
57 E W R F F E C P K C H Y R F T T Y V G N K E I E N L I R
25592 tttagaataacttgcgcagctaaatgaagcaggaaacttcaaaaaggagctgctgctaatcaaaacatttaccattcatatcga
85 F R N T C R A K M K Q E L Q K G A A A N Q N T Y H S Y
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25760 gaatgggtatctatatag 25777
141 E W V S I *

dp1ORF046

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42942 gggattgacaaaacgacagtagcaatcaatcaccaaaatgacgtcattcaagacggaactagaaaaattcaacgttaccgtctt
57 G I D Q T T V A I N H Q N D V I Q D G T R K I Q R Y R L
43026 tatcagcacttaaaaagggaagtataacaggctatacaactctcgaccatttttagagagctctctattttatcgaaggttat
85 Y H D L K R E V I T G Y T T L D H F R E L S I L F E S Y
43110 aagaaccttggcggaaatgggtgaagcttgaagccttgaatgaaaaatacaagaattaccaattagggaggaagatttagatgaa
113 K N L G G N G E V E A L Y E K Y K K L P I R E E D L D E
43194 actatctaa 43202
141 T I *

dp1ORF047

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47626 caagtcaaaagtcttctgctgctctaaaagagtagatgaaagaaaatgacattgaatctgctcaaggtaagcacttttctgct
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47710 accttctacacgacagagcgtcactatggcgaagaagcgttgaagaattatcgaaaaattagttgacgaagccgagacg
57 T P Y T T E R S T M D E A R L K E I I E K L V D E A E T
47794 gaagaaatgtgtgaaaaactttcagggttatcgaaatacaagcctgtcatcaatcgaacttctcgaggatagatttatcac
85 E E M C E K L S G L I E Y K P V I N T K L L E D M I Y H
47878 ggcgagattgaccaagaagcaattcttcagcagctgtcatttctgttacagaaggcattcgttttggaaaggctaaaatttag
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113 G E I D Q E A I L P A V V I S V T E G I R F G K A K I *

dp1ORF048

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29 P V K E T E K Q Y K V T G I N P N L Y L D L G S V I R K
16541 agcgaacttgacattgcagttatcaagcagctcctgtcgctgaaactggagtcacacttactcgcgacatggaagttgatgct
57 S E L D I A V F K A C P V A E T G V T L T R D M E V D A
16457 agaattgaaatcatcaagaaatttaactacaagaatcgaacgccttaacgaaagaattaaagcaagaaatgaacaaggtaaaaca
85 R I E I I K K L T T R I E R L N E R I K A R N E Q G K Q
16373 gaaagccgccacttagtatctgcgctagaagattgcgctcgtcaaattgctggaatttatcaataa 16308
113 E S R H L V S A L E D C A R Q I A G I Y Q *

dp1ORF049

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43850 agcatttgcgttagtcaagcgatagacgttagttagttagcttgcattgacacacgctcactcgtggtcgttgacggaat
57 S I C V S Q A I D V V V R L T C I V P T L I V V D G N
43766 tccgtcgtaggcgttagtgcagtgatgttatcactgtcaatgaacatccctgtatgacctccagcgctcgtcgttagcacc
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dp1ORF050

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379

15165 cttcttgctgtaaaggaccagcggaagaactcgaaacaccttgagccctttgtgggatacattgacaatctagtcgaatgtttt
29 L L A V K D H G E E L E N L E A F V G Y I D N L V E C F
15249 cctgaaagccaacgaatgtcttgaggctatgtgtattagatgaccttccagtcactaatcgccgctgaaattggataccac
57 P E S Q R N V L R L C V L D D L P V T N A A A E I G Y H
15333 tatacatgggttcaccaacttcgagacaaagcagttgaaacacttgaagaaatttttagatggggataacattattcgctctaaa
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29 S W V S D G Y G G K K K D K A N E V V A D D L V C L V D
29933 aattcaactgttctgaccttttagccaattctactgacgggaaaaatttttgcccaaatggagtgaaaaattttcattcta
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30017 tatgatgaaggcaaaatcattcaacgagccgatactatcgaaattaaaaactcaggaagacgggtacagggtagtagaaacccac
85 Y D E G K I I Q R A D T I E I K N S G R R Y R V V E T H
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dp1ORF052
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29 V E F D E Q D T D R P D D Y I V L R Y S H R M P S A T N
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30768 gttcgaacattatcaaggacatgggtacgaagtaacctatcgagaaactgggtgactactcgacacaaatgctttctagatagc
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30852 cgactagaatcgaatatagaattccacaaggaggaaactaa 30893
113 R L E I E Y R I P Q G G N *

dp1ORF053
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29 P F F Q A V A S I L S I V H D L P C P G R A I M T I K S
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113 S S F G I I F A I A M L L S T *

dp1ORF054
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57 V M A L P V S H A E D L G K R L C I A N S R L E A F R E
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113 G N L A L Q L V E S G A L *

dp1ORF055
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29 A A Q I P A T A A T Q V G N K K Y I L A G T C V K N A T
27795 acatttgaaggacgcaaaactggactcgaagtagtatctaccggtgaacaattcgacggagttatcttctgctgaccaagaagtg
57 T F E G R K T G L E V V S T G E Q F D G V I F A D Q E V
27879 tttgaagggtgaagaaaaagtaaccgtgacagttatagttcacggattcgtcaaatatgcagcccttcgaaaagttggcgatgct
85 F E G E E K V T V T V L V H G F V K Y A A L R K V G D A
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113 V P E S K N A M I L V V K *

dp1ORF056
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29 V P G T P Y R L Q V W V K M S L V K I E T R A G N G Y Y

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85 K S L A E Y C E P M N R H I L E T I A S R E A A E L N R
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113 A K K Q D Q Q K W R Y *

dp1ORF057

380

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10195 gcagaaaggcttttaagaaaatga 10218
113 A E R L L R K *
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29 K V Q P N S G A T D Y Y K G D V V T D S M L I E C K T V
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113 R N D N L I *
dp1ORF059
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29 I T V C E V A A T K M E E Y A K T H A I W T G N A
30322 cgacagaaactcaaaggagaagctgcttgggtaagcgcagaccaaatcatgatagctgtatcacatcacatggactacgggtt
57 R Q K L K G E A A W V S A D Q I M I A V S H H M D Y G F
30406 tggctagaactagctcatggtcgaaaatacaaaattctcgaaacggctgtagaagacaatgtcgaaagacttttagagcgtg
85 W L E L A H G R K Y K I L E Q A V E D N V E E L F R A L
30490 agaagggttattagactag 30507
113 R R L L D *
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29 A S P L G P S S R I H V K S S G T N S L G F L L V L R T
37902 ccaatgtatttcccagattctgcattaaaattagtccttaaatgtcatctcgctatctaataacaacttgggactcatttaca
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37734 tttgaaaggctgctgtag 37717
113 F E R S S *
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29 F D E E A T Y D R Y R E A L E K V G N V A Y F C E I D T
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57 G N L V I E L E L D S L D D L I A L S N V V G T G L K L
19223 tcacggccttatagagaagataagcctttcaattatggattgttgacgggtacatggaataa 19161
85 S R P Y R E D K P F Q L W I V D G Y M E *
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29 R C R S N R C K K F L L V F C Q P F C A N S N R N T F S
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85 F N N S T P K S L S N R H H A F F R S R F S N S R F L
44948 actaaactga 44940
113 T N *
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47200 atgaaattcactgaaggaaaaaattgggtataaagttggagagatatgtcaaatgttgaaacgctctctatctacgattaatggt
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29 W Y E A K D F A E E N N I H F P F V L P E P R T D L D H
47368 cgtgggtctcgtattctgggatgacgaaggcgtgaacaaactcaaacgatttgggacaacctaatgcgcgggtgacttggcattc
57 R G S R F W D D E G V N A K L K R F R D N L M R G D L A F
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47536 aattaa 47541
113 N *
dp1ORF064

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29 acaattgaatcagttgaagaaattgacgaagttgaacaaatgcgcgaagagtagtcggcgttaaaccgttcctgagctcgttgaa
57 T I E S V E E I D E V E Q M R E E Y A A K T V P E L V E
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29444 gagtaa 29449
113 E *
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57 A L P N Y F A R C S K I P F Q P L V S I E P S I V S T *
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29 F V S V S E L S N F L R V D S D L K T C F F S D E F L S
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85 L V I S V S V Q D H S S R A N T C T I F D V I H C C *
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29 G A G F R T V D F S L T E P T S S G C S L T S G I S S S
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85 A A S S F L G S V S S S I V Y Q R S R V E A E R *
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29 S A L L D K H K S V A Y V S Y M I C L M K T R N D V V T
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29703 cttgaaactgatgaaaagtcgaacgctggttcgacaatcttaataagaaagggctgatgggacatga 29768
85 L E T D E K S N A G S T I L M K R A D G T *
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20411 atgaaactttatcacgccactgattttgataatcttggttaaaattctagctgaaggattgaagccttcagctggagtattttac
1 M K L Y H A T D F D N L G K I L A E G L K P S A G V I Y
20327 ctacgaaaagttatgaaaaggctctagcctttttatcgcttcgaaatgttgataactattgtcgttctcgaaacttgaaatagat
29 L A E S Y E K A L A F L S L R N V D T I V V L E L E V D
20243 attgaaaaatgtactgaaagtttcgaccataatgaaaagatgtttgttagcctatttcttcgacactgtcgcgcttgagat
57 I E K C T E S F D H N E K M F C S L F H F D T C R A W T
20159 tatgacaagacaattgaagtagacgacattgacttttcgaaagctcgaaaatgatagaaagtga 20094
85 Y D K T I E V D D I D F S K A R K Y D R K *
dp1ORF070
15973 atgataaccttattttaaaataaacagtgaaaggaacagttactccaattaaagggctcagccatgcaactgtacgcagaccttatt
1 M I T L F K I N S E G T V T P I K G S A M Q L Y A D L I
16057 cctatacaagaggacgatatacagttcgttgatataactggacttgacctattgttcgagaaaacgtacttgagctcatttca
29 P I Q E D D I Q F V D I T G L D P I V R E N V L E L I S
16141 cggagccgtgtagaggtttcaaatatgggtacaaacctcgaccagaatgatgtcgacgatttctacagcagcgaagaagaa
57 R S R V G V S K Y G T N L D Q N D V D D F L Q H A K E E
16225 gcgctcgactttgtaactacctaaccaagctacaaagtcacaaaagcgaataaatag 16284
85 A L D F A N Y L T K L Q S Q Q K Q N K *
dp1ORF071
38904 gtgaaacaggtcctagaggagttcaaggtcttcaaggtcctcaagggcttcaaggaattcctgacctgcaggagctgacggac
1 V K Q V L E E F K V F K V L K G F K E F L D L Q E L T D
38988 gttcgaataatactcacctcgttttcttaataatagtcacaaacggtaggggatttagtcatactgacagcggagcagcatagctcg
29 V R N I L T S L S L I V Q T V R D L V I L T A D E H T S
39072 gtcagtatcaagatttcaatcccgctccattcaaaacccctgcagcctatacatggacgaatggaaggggaatgacggagctc
57 V S I K I S I P S I Q K T L Q P I H G R N G M T E L
39156 aagggatacccggaagccaggcgacagcggaagactaattttccatatag 39209
85 K G Y P G S Q A Q T V R L I I S I *
dp1ORF072
51045 atgttctcttcgcttcaagttgtctcgaaagtttttcaattatttgttcaggagtcgcttcaatttgaagaccatttactttca

382

1 M F L R L Q V V S K V F Q L F V Q E S L Q F E D H L L S
50961 tcaaaatgcttcaactccttcccttgtaaccttacttcgaagacgagcagtcgacctagaggcttttgccttcaatggagagct
29 S K C F N S F P C N L T S K T S S R P R G F C F Q W R A
50877 ttgcctttttcagttccttcttcgcttctcttgaatcctataagagtatagggtccagtttcaacgtcccatatattc
57 F A F F S S F F A F L F E S Y K S I G S S F N V P H I F
50793 gatgatttttcggtcttcgcatatcggtttttaacgacagatag 50749
85 D D F S V F A I S V F N D R *

dp1ORF073
14262 gtgaacgcttgccggaagaatacgcgaagaacttggaacctatcactgaagcagaatacatcaagcgaacagaaaaaccta
1 V N A C R K N T T K K L G N L S L K Q N T S S E Q K N L
14346 aagcagttgcaaaacctactcgaaaaactccagcgcttctcgtcgccctcgcccttaaaagaaagggtgaataaaatgtgtg
29 K Q L Q N L L E K L Q R L L V A L A L K R K V E I K C V
14430 aaaattgtcaaaacgaaacattcaataactagattttcaatgaagatgaaagtggctatgtcgacgcctcattcactacaagg
57 K I V K T K H S I L E F S M K M K V A M S T P H S L T R
14514 agattcgacacccgcagcagctatttagcaatcgagcggtag 14555
85 R F A T P Q Q L L A I E R *

dp1ORF074
32298 gtgacgaaaaagaaaatccaggattgcaaatgcttatggagtactattttcagtcgctcctcttttgcctatagaaggaaa
1 V T K R K I Q D C K C L W S D Y F Q S L L F L Y I E R K
32382 ttacatggatttttgggtcaattgcagcaaaaatgactttgggatactcctcaaaactcacaagtcaattaaatcttgctcaaaagtc
29 L H G F W V N C S K N D F G Y L K L H K S I K S C S K S
32466 agcgcaacggctcgactagagctcttcgaagtcctttcaaatgggtctgttttaacaggattagggaaggacttacgactgc
57 S A T A R T R V F E V L S N W F C F N R I R E R T Y D C
32550 ggttacccttctcttatgggatttgcagcgcctctattaa 32591
85 G Y P S S Y G I C S R L Y *

dp1ORF075
22447 atggcagaagtttttgcctgtgaattccgtcatggcccaagggaagaatgaaagagccatcgatactgttttctgaacgaatg
1 M A K F C P L N S V M A Q R E N E R A I D T V F P E R M
22363 gaacgctctgctatgacgatatcgaaagttcgaaaagggtgagccctttgtccaccatgttaggagctggagttgtttcttacta
29 E P S A M T I S K V R K G E P F V H H V R S W S C F L L
22279 aaagggaacgaagtgaacttaggttagtttatttctcagccttattgtcattatcagtcactcctttaatgtaggaaacctgttgc
57 K G T K L N L G S L F L R L I V I I S H S F N V G T C C
22195 gtcactaaattcttgcacaaaggcttgagctgcttattag 22154
85 V T K F L P N G L S C F I *

dp1ORF076
5728 gtgagagcatttttcttactcacgtcttcgagcaagtggctgaatgtagggtactcttcatcttctgtaacaatatcaatattg
1 V R A F S S L T S S S K W S N V G Y S S S S V T I S I L
5644 tactaccattcccaataacttttagcgaagatttccaggaactaatgtgacggttgcggccggtgcttttctacaagttt
29 Y S P P I T F S E D S S G T N V T V A A V V F S T S F
5560 ccaaaactgctctgctttcacaaatcagtcattttcaacactcgctgtcgataatgcacgaaggaagttgagccatcacgct
57 P N C S A F T I T S I S T S L S I M H R R K F E P S Y A
5476 gtaaacatgacgcattcgccgtcaccaaaaatagccaatag 5435
85 V N M T H S P S P K I C Q *

dp1ORF077
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1 M E R I K T L F H V I Y A N G T H L E V A A L F D T V D
14884 gattatgatgacgttatagaggacatccaggggtatattgataccctgacctttataatcaaaggagcattagaatggcgctt
29 D Y D D V I E D I Q G Y I D T P D L Y N Q R S I R M A P
14968 tacaatcctgacatcaatggtagcgtattgctactgacattttactacgactagatgatattatctacgtcgacgcaactgt
57 Y N P D I N G D A I A T D I L L R L D D I I Y V D A T C
15052 gaaactattaaatcagaggagcctattgcatga 15084
85 E T I K Y E E P I A *

dp1ORF078
17507 atggcaacagtaaaaggaaacagtaaaatttgacggacgtcttgaactatcttcgactacgacgatttagagtggaaggat
1 M A T V K E T V K F D G R L V T I F D Y D D L E W E G Y
17423 gcacctaataaggattcgaaagattgtaggacatggaagtccttagcattcgagttcgaaacgaaggtgaggacgacgagtg
29 A P N E G F E D V E D M E V L S I R V R N E G E D D E W
17339 gttgaagttatcgctgctatgaaaacgatgacgaggacgaagatttgaagggttataa 17280
57 V E V I A C Y E N D D E D E D L E G L *

dp1ORF079
35288 atggaactgataccattgataaatcctcgaacaagggtgacccctgcgcttaccatttgcagcgaatccagtaaccttagaa
1 M E L I P L I N P R T R L T P A L T I C P A N P V T L E
35204 acaattgaagttcccatgctgcaatttttagagacagctgaaccaatcattgacccaataccactaatgaagtttcaatcagg
29 T I E V P M L P I L E T A E P I I D P I P L M K F R I R
35120 ttgcacctcctgaaacctctgtcccacaaagctagcaatcttgtaactaatgatgaaagcatgtttccagctgtcgataaa
57 F A P P E T I C P T K L A I L L T N D E S M F P A V D K
35036 agtgagcggagaagtgaagcaataacctga 35007
85 S E P R S E A I P *

dp1ORF080
42490 atgttgaaccttacaataatcgcgcaaaatttggtggcagagttcactattggacaaggagctgaaaagaacttgcctaaacaacg
1 M L N L T K S R Q I V A E F T I G Q G A E K K L V K T T
42574 attgtgaacattgatgcaaacgcagatcaacgcgtctctgaaactcttcatgaccacgactgtatgctgcgaacgcgtcgagaa
29 I V N I D A N A V S T V S E T L H D P D L Y A A N R R E
42658 cttcgagctgacgagcaaaacttcgcaaaactcgtaacgcaatcgaagatgaaattctagctgaacagctcaagactgaacaa
57 L R A D E Q K L R E T R Y A I E D E I L A E Q S K T E T
42742 gctctaacagctgaataa 42759

dp1ORF091
43189 atgaaactatctaacgaacaatatgacgtagcaagaacgtggtaaccgtagtcgttccagcagcgattgcactaattacaggt
1 M K L S N E Q Y D V A K N V V T V V V P A A I A L I T G
43273 cttggagcgttgatcaatttgacactactgctatcacaggaaccattgcacttcttgcaacttttcaggtactgttcttagga
29 L G A L Y Q F D T T A I T G T I A L L A T F A G T V L G
43357 gtttctagccgaaactacaaaaggaaacgaagctcaaaacaatgaggtggaataa 43413
57 V S S R N Y Q K E Q E A Q N N E V E *

dp1ORF092
46989 atgaaaactatctccatattaaggaaagacactaaaaggagccggacaggaacggaagaaaaactgcactcgaactagctcaa
1 M K T I S I L R K D T K R K P D R N G R K T A L E L A Q
47073 gagattgatattgtcacctagttagcttagcagagctccttcaaattcctgaaaggacggcaaccagaattttaaaactcgacaaa
29 E I D M S P S E L A E L L Q I P E R T A T R I L K L D K
47157 ctgctcaacaaagagcaatgctcaataatagaaaggtatataaatgaaattcactga 47213
57 L L N K E Q C S I I E R Y I N E I H *

dp1ORF093
45756 atgcaacatacgattaaacaatgtttgaaacttgcttctgctgaactgcaatatcaattgctgttttagtttccctaaacct
1 M Q H T I K Q C L K L A F L L T A I S I A C L V F P K P
45672 tgctcatcgctaaaaggaaacatggatgcttctgtgctattcgaaacattcaacctgggtgcgcgaatggagtagtcttgaa
29 C S S P K R K H G C S C A Y S K H S T W C A N G V V L N
45588 gaaaactgctcattgcttgaagaagctattcggtttcgagagtcattgtag 45538
57 E N C S L L E E A I R F R E S M *

dp1ORF094
8281 atgtacgaattagtcttatcactaaaattgacgcgcagacgcgcgatgtctcaagatgtcgaaaagtgttcaaaaggctcaag
1 M Y E L V L S L K L T P T A P M S Q D V E K C F K R L K

8365 tatattcagtgccgcgcaggtgaatgcattaaaattgcacacggatttgccttgaacttcctaagggatagaagcaattcttgc
29 Y I Q W R Q V N A L K L H T D L L L N F L R D M K Q S C
8449 atcctcgctccagctcttttaagaaaactggcttaa 8484
57 I L V P V F L R K L V *

dp1ORF095
8877 gtgggaaaactacttcagctctcgacattgtcaagaatgcgcaaatgggtatttgagcaggaatgggaacagaagactgaagaac
1 V G K L L Q L S T L S R M R K W Y L S R N G N R R L K N
8961 tcaaggaaaagctggaaaatgcgcgtgcatccaaagctagcaagactgctgtcaaggaaactgaaatgcaactcgatagcttcc
29 S R K S W K M R V H P K L A R L L S R N L K C N S I V F
9045 aagagcctcttaagattgtatatcttgaccttgagaatacattag 9089
57 K S L L R L Y I L T L R I H *

dp1ORF096
46681 gtgattcataaattcttcaatttcgttgaacttatctcggttttctctgttaccaggttgcaatttgactgtcttcgaaagtat
1 V I H K F F N F V E L I C G F S C Y Q V A F D C L R K Y
46597 ctttagcaagaggttcaataaccttttcccaattgtaaatatcacgcaggactttcttctgtggatacatctctcgacaatttc
29 L S K R F N N L F P I A K Y H A G L S L L D T F L D N F
46513 gatacatcttctgaacttgcaagacttgacatcttgagtagttaa 46469
57 D T S F E L A R L D I L S S *

dp1ORF097
39100 atggacgggattgaaatcttgatactgacgcagctatgctcgtcgctgctcagtagtaaatccctccacggttggactatt
1 M D G I E I L I L T D V C S S A V S M T K S L T V W T I
39016 agagaaaagcaggtgagtagatattcggaacgtccgtcagctcctgcaggtccaggaattccttgaaagcccttgaggacctggaag
29 R E S E V S I L R T S V S S C R S R N S L K P L R T L K
38932 acctgaaactcctcaggacgtgtttcacctatcttggaaactga 38888
57 T L N S S R T C F T Y L G N *

dp1ORF098
43627 gtgaaaatgctccgtgggatgctaaacgagcgacatcttcatctggggacgcaagggtgctagcgcaggcgtggaggtcata
1 V K M L R G M L N E A T S S S G D A K V L A Q A L E V I
43711 cagggatgttcattgacagtataacatcattcactgcaactacgcctacgacgaatttccgtcaacgaccacgatgagcgtt
29 Q G C S L T V I T S F T A T T P T T E F P S T T T M S V
43795 ggtactatgcaggtcaaccttactactacgtctatcgttga 43836
57 G T M Q V N L T T T S I A *

dp1ORF099
38298 atgcaagttcgccatctgctactgaagctccagctgggtggtggtctacgcaagttcctaccgtcccaggtggtcagttattat
1 M Q V R H L L L K L Q L V D G L R K F L P S Q V V S I Y
38382 ggactcgaacaagatggcgctacactgacaaactgatgaaattggatattcagtttcaagaatggcgagcaggggtcctaaag
29 G L E Q D G A T L T K L M K L D I Q F Q E W A S R V L K
38466 gtgacgcaggtcgtgacggtattgcaggaagaacggaatag 38507
57 V T Q V V T V L Q E R T E *

dp1ORF100
1597 atgcagttgacaccaagcgagttctatttggatttagaactacggctgagaatatgtcaagattccttacctggactctcacgg
1 M Q L T P S E F Y L D L E L R L R I C Q D S L P G L S R
1681 agcttatgtggaagcatgctcgtatcgactctataaactatgggaaactcctacaggttgcgcagaatgtacttactacgaga
29 S L C G S M L V S T L S N Y G K L L Q V A Q N V L T T R
1765 ttttcacagaagcagattgaaatgttcaagaacgtaa 1803
57 F S Q K T R L K C S R T *

dp1ORF101
19220 gtgataatttttagtccagttccactacatttgaaagcgcgattaggtcatctaggctgtctagctcaggttcgattacaaggt
1 V I I L V Q F P L H L K A R L G H L G C L A R V R L Q G
19304 tgccagatcaatttcacaaaagtaagcgacatttccaaacttctctagtgttcacgatacctatcatatgtcgcctctcgt
29 C Q Y Q F H K S K R H F Q L S L V L H D T Y H M S P L R

385

19388 caaatagtcgagcagaataaacttcgaatttcatttag 19426
57 Q I V A Q N K L R I S F *
dp1ORF102
4034 atgataacgtgggaatgtttgactgtatcgccgaactcgataaaattcctgggtgatttagacagcctaagacacgtgaacagc
1 M I T W E C L T V S P N S I K F L V Y L D S L R H V N S
4118 ttttggaaagcaccacaaatttcttgggataattatctatacatgcgcgagcgaatgggttgagaaaagacaagctcttacctattt
29 F W K H H K F L G I I Y T C A S E W L R K T S S Y L F
4202 tccatattgggagaagactttaaatggctcaactga 4237
57 S I W E K T L N G S T *
dp1ORF103
49352 ttgaatcatagatatagtaaacatcacaaactattttctttggcagattgtctttctttgtatttgcgcgggtgtcctattgt
1 L N H R Y S N I T T I F L W Q I V F L C I C C A V S Y C
49436 gcaggagtgcataatgagcgaagctctcaagataaggtgattcgaagtataagcagaaaagaaagtcagcgtctacttgaca
29 A G V H N E R E S Q D K V I Q S Y K Q K E K S A V Y L T
49520 gtcgatagttcaggagcttggctaggaagtgtccgggagccaaggaaagtcctctacatgaaaaggacagcatgttagga
57 V D S S G A W L G S A P G A K E S P L Y N E K G Q H V G
49604 aaattgaaaagggtgggagagtga 49627
85 K L K E V G E *
dp1ORF104
21427 atgagaaaaagagtgtatttgaagctaaaaagggtgaactggatgtccttaattcctactctcgaatgggtgagtttttcgaa
1 M R K R V I L K L K R L N W Y V L N S Y S R M V E F F E
21343 cttttgaacttttcgaatgggtcgaacttttcgaaggattgaggttttcgaaccgggtgagtttttcgagcattctcgacttttc
29 L L N F S N G S T F R R I E V F E P V E F F E H S R L F
21259 gaccctttctatgtctcgacttttcgagtggtttga 21224
57 D P F L C S T F R V F *
dp1ORF105
2028 atgatagtcgcatccaccagttcgaatgaaaatagctttttgacctataaccattccttcaccttgattgtaggaccgaaaat
1 M I V A S T S S N E N S L L T Y N H S F T L N C R T E N
1944 ttccatgataggcattttctcagggtcggaacattgattcgaatccttgcctccttcagggtgattgattgattaccattat
29 F H D R H F L R V A N I D S N L A S F R L I V L I N H Y
1860 cctgtcctgtctctaaaatttcgaggacagtaa 1828
57 P A P A L K F R G Q *
dp1ORF106
10529 atgaacctcgatcaatgatgtaaaactttgaactcgctgtccatagacttgatctagaatcttcaataatgtttcgaacattttc
1 M N L V N D V N F E L A V H R L V S R I F N N V S N I F
10445 tacccattattagaagcagcatcaatttcaataggagagccaagtcctttgttcacatccttcgagaaaattcgagcagtagt
29 Y P I I R S S I N F N R R A K S F V H I L R E N S S S S
10361 gggtttaccagttccagcgcaccacagaatag 10329
57 G F T S S S A T T E *
dp1ORF107
10750 atgagcgtgacgcccctttcgtttattgggaaacttgcaaatggaggaatgcgtgacagtatcacaaggctcgaaaaagtccttg
1 M S V T P F R L L G N L Q M E E C V T V S Q G S K K S L
10834 attatagtcacacgttgacatggaagccgtttcctaagtcactag 10878
29 I I V I T L T W K P F L M H *
dp1ORF108
49447 atgcactcctgcacaataggacaccgcgcagcaatacaagaagacaatctgccaaagaaaaatagttgtgatgttactata
1 M H S C T I G H R A A N T K K D N L P K K N S C D V T I
49363 tctatgattcaatttcgcttacctccaatcctcttaccattgcttgctgaaatctagaaccactgaagtatcatatatacga
29 S M I Q P R L P P I L L H C L P E N L E P L K Y H I Y D
49279 tataaagcctttggcctaaaagggtcaataa 49250
57 Y K A F G L K G Q *
dp1ORF109
31632 atgtgggtgtcgaagtcctcaaatagttgatttctccttcaactttccagcctttgaaagccttacctgttaaggtagggtcaact
1 M W L S K S Q I V D S P S T F Q P L K A L P V K V G S T
31548 gggtttgggaaaattcttctacctgttcaactcgaactgcgtcgccggttccgttccaccgttcaatcgaaatgtcacgcga
29 G F G E I F L P A S T R T A S A V P V P P F K S N V T R
31464 cgaagaaccgctggaagtgtgccacatag 31435
57 R R T A G S C A T *
dp1ORF110
16444 atgatttcaattctagcatcaacttccatgtcgcgagtaagtgtgactccagtttcagcgacaggacatgctttgaatactgca
1 M I S I L A S T S M S R V S V T P V S A T G H A L N T A
16528 atgtcaagttcgctcttcttaataactgagcctaggtctcaagtaagaattgaggtgattccagtgaccttatattgtttctca
29 M S S S L F L I T E P R S K Y K L G L I P V T L Y C F S
16612 gtttctttacaggaatgctttcatag 16638
57 V S F T G M L S *
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28657 gtgactctatcaagaagctcttgcaattgggtgtcaaggttcttgggaaaacttcttgcttcttgcaagtgacgctgagaaat
1 V T L S R K L L Q L V F K V L G K T S C F L Q V T L R N
28741 tcacgcgtgaaaaacaggtcttcaaatcgctgtctactctcaagaaattgctcagttcgctgacgctgacaaacttctgacg
29 S S L K K Q V F K S L S T L R K L L S S L T L T N P L T
28825 ttggtaacattcgctcagttcaactga 28851
57 L V T F V S S T *
dp1ORF112
32207 atgcaaaactgatttaggcaaatactgcttcgacgcagcagccgttgcttatattagatatttgcaggaagacaagactcctagg
1 M Q T D L G K Y C F D A A A V A Y I R Y L Q E D K T P R
32291 tatcctgggtgacgaaaaaataatccaggattgcaaatgcttatggagtga 32341

386

29 Y P G D E K K N P G L Q M L M E *
dp1ORF113
17715 atgaaaacaggttaaagaagcaatcaacaattcggatgaatgggtacgaaattatcaacgaaaacggccaatgattcaa
1 M K T V K E A I K Q F G D E W W Y E I I N E N G Q M I Q
17631 gacggaagaatcgaagacatggcggaatacatggaagaacggcgaccaagttaagttcatcaactatggtagacatcgaatct
29 D G R I E D M G E Y M E E T V D Q V K F I N Y G D I E S

17547 caaattatcaaactatatatcgcataa 17521
57 Q I I K L Y I A *
dp1ORF114
52952 atgctattggcgaagacggggaaacagtcctcctgataattgtccattatgccaaaacggattccctcgattgaaaaactat
1 M L L A K T G K Q S I L I I V H Y A K T D S L V L K N Y
53036 ttcttcaactttacaaccatgatacgggaaaagtgaacatgggaccgagccgttcttatgttcaaaagattgttacattta
29 F P N F T T M I R E K L K H G T E A V L M F K R L L H L
53120 tcaataaatatggaagccttgtga 53143
57 S I N M E A L *
dp1ORF115
5342 atgagcctccttttttgatatataataatacacgaattatcgcgagtttgtaaagccgtttctaaataattttaaatctttt
1 M S L L F L I Y I I Y T N Y R E F V K P F L F K S F
5258 aagcatattgagttttgcttcataagtcctgttcacggcagcctcttgcatattgagtacaatgaaaggaggttcctcgatatt
29 K H I E F C F I S P V H G S L L H F E Y N E R R F L D I
5174 gttgaaactatagaaggtgaataa 5151
57 V E T I E G E *
dp1ORF116
20662 atgaaattttcaaactttgctaaagcacttactaatgaatacctaaggttagtgaacaatgaccaagctgaagctcttagggcga
1 M K F S N F A K A L T N E Y L M V V N N D Q A E V L G A
20578 ggaaatatcgaaaacattctcaacgggttcgaactttgctaattgttagctgaagcgacagttttaaactcgaaaaactcagc
29 G N I E N I L N G S N F A N V V A E A T V L K L E K L S
20494 gaagaggaagctattgagtag 20474
57 E E E A I E *
dp1ORF117
24680 atgataacaggctgctcgaaacttttaaatcgaagtgaatctcgtaagtcactaatagttttgttcaagttatctgctactgtg
1 M I T G C S N I L N R S E S R K S L I V L F K L S A T V
24596 ataaggctcttgacatcgcttgccttatatgtcatttagtcaatgggttcattaagaataaactcgacaaggaatttgcctcaag
29 I R S L T S L V P Y M S L V N G S L R I T R Q G I C F K
24512 ccggttggggcggtattcttga 24492
57 P V G A D S *
dp1ORF118
15023 atgatattatctacgtcgacgcaacttgtgaaactattaaatacaggagcctattgcatgaacaatcagcgaaaagcaaatgaa
1 M I L S T S T Q L V K L L N T R S L L H E Q S A K A N E
15107 caaacgaatcgctgaacttcggaagactatcaacgtgcaagaggtcgaaataaacttccttctgctgtaaggaccacggcga
29 Q T N R R T S R R L S T C K R S N K L P S C C K G P R R
15191 agaactcgaaaaccttga 15208
57 R T R K P *
dp1ORF119
41054 atggagggttcaacatccccgattcagtagctcctacttttccgggcatttcttttagtagacagacttcagcggttcgacagat
1 M E V Q H P R F S T S Y F F G H F F S R H D F S G S T D
41138 tttaacagggaacaacttctccaaatcatgtcgaaacttcaagtcacttcaacaatgcttcggcgcttaccgacccat
29 F N R E Q L P P N H V E H S S Q L Q Q C F R R L R I H Y
41222 ccaagcatttcacgctga 41239
57 P S I S R *
dp1ORF120
28387 gtgttgaagcgcaagcagaatacatcggtatgcaattgcttcaatacggtaaatcactgtcaaatcaactaacagcgaggtc
1 V L K R K Q N T C V C N C F N T V N S L S N Q L T A R L
28471 aatacacttacgactacaacatggatgctgaagcaaatatgcagtcactaagaaatggactaacccagctgaaagtgaacccta
29 N T L T T T T W M L S N N M Q S L R N G L T Q L K V T L
28555 tcgctgacatttttag 28569
57 S L T F *
dp1ORF121
39222 gtgcagacggatcacgtgagttcagtttggagaataataatcaacaatatatgggttattactccgattatgagcaagcagata
1 V Q T D H V S S V W K I I I N N I W V I T P I M S K Q I
39306 gcagggtacgaactaagtatcgatgggttggaccgccttgccaatgttcaagtgagggtcgaaaacgagttccttaattctttat
29 A G I E L S I D G L T A L P M F K W E V E T S S L I L Y
39390 ttgaatttgggttaa 39404
57 L N L V *
dp1ORF122
40402 atgttattctccttatcctacatacgaatcacgttcatgtctggttaaacgagattgttccggttctaatacggccgacttg
1 M L F S L S Y I P N H V H V W I K R V L F R S K S A D L
40318 aatggattgggtaaagatcccgttatcgatgtgaatgaaccttgcgtaaggtacataacttcattccctcgaggagaacataga
29 N G L G K D P V I D V N E P L R K V H N F I P C G _ E - H R
40234 aattcgggtcacttga 40220
57 N S V T *

dp1ORF123
21327 atgggtcgacttttcgaaggattgaggttttcgaaccgggtgagtttttcgagcattctcgacttttcgacccctttctatgct

387

1 M V R L F E G L R F S N R L S P S S I L D F S T P F Y A
21243 cgacttttcgagtggttttgaggttttcgagcaggttcgacttttcgagaaattgagtttttcgacctctaaattaggctcgatt
29 R L F E C F E V F E Q V R L F E K L S F S T S K L G S I
21159 attcgaaaagtttag 21145
57 I R K V *
dp1ORF124
17891 atggtaaaagttaaagatttgcaagtaggaatgaaagttgtaaatgcaaaaggtagtgaatttaaagtaactgaccgtcaagggt
1 M V K V K D L Q V G M K V V N A K G T E F K V T D R Q G
17807 cgtaaatgggtaagcctagaacgtcttagtgatggacgtattcggttctatgataacgaatcactaatggacgaaaaagtgag
29 R K W V S L E R L S D G R I R F Y D N E S L M D E K V E
17723 gtagtaaaatga 17712
57 V V K *
dp1ORF125
49916 atgtcctcagccgcttcggttaaaattggaacaagtgaattatatagatgctcctcttttagcttgctgataagggtattcatca
1 M S S A A S V K I G T S E L Y R C S S F S L S I R Y S S
49832 gtttcgccatttcgaaaaattcgaaatccaggaaatgggtcgagaatagtttcgctcgtccggaactcttccatattcgcgaaaag
29 V S P I S K N S N P G K W S R I V S S S G T L P Y L E K
49748 tgttcttga 49740
57 C S *
dp1ORF126
16136 atgagctcaagtagcttttctcgaacaatagggtcaagtcagttatatcaacgaactgtatatcgctcctcttgtaggaata
1 M S S S T F S R T I G S S P V I S T N C I S S S C I G I
16052 aggtctcgtacagttgcatggctgaccttttaattggagtaactgttctcactggtttatttttaataaggttatcatttct
29 R S A Y S C M A D P L I G V T V P S L P I L N K V I I S
15968 atcctctaa 15960
57 I L *
dp1ORF127
13511 atgctaaatagctttccattcaccgtcgctgttcttgcgccatttttcagtttcacgatactgaccaactttgcaaaggctcgt
1 M L N S F P I H R R C S C A I F Q F H D T D Q L C K G R
13427 gaaatagtgctacgattgcaactgtttccattgggttaaatgtcttccagcctttgacctaccatgggtatccatttcgaaaagta
29 E I V L R L Q L F P L G K C L P S L C L P W Y P F R K V
13343 gttgattga 13335
57 V D *
dp1ORF128
4852 atgacagcagttcaacaagtttaagttctacttagaagaagccgctcactttctaaaagatgttgagtacagtgacaactta
1 T A V Q Q V K F Y L E E A G A H F L K D V E Y S D N L
4936 gagcaagcaattatgaaagatattcttaaatggaatggcgctcatagagatgagcacgatatgaaataaacttcatacgaagta
29 E Q A I M K D I L K W N G A H R D E H D M K I T S Y E V
5020 ttatag 5025
57 L *
dp1ORF129
25133 atgaactttctgctaaagcaacttgctcactggaagttcaactaatgtacgcagccaccaatcttacattgaagaattcagta
1 M N F L L S N L R S L K F K L M Y A A T N L T L K N S V
25217 agaaggaaaaagcggaaggaatgggaacgattttggaagaacttgctcagcttgacgaaatctcagctggagcattgcctg
29 R R K R R T R N G N A F W K N L L S L T K S Q L E H C L
25301 tattag 25306
57 Y *
dp1ORF130
16789 gtgcttgactttattcctttattatcgatataatcataatataaataaaaacagcgctcaaggacgcagaaaagggtcaattatgg
1 V L D F I P L L S Y N H N I N K T S V K D A E R G Q L W
16705 aaacaacactttatttcggttatcttacagcagattggaaagacggtcacagaactacactttccactatgaagcattcctg
29 K Q H F I S V I L Q Q I G K T V T R T T L S T M K A F L
16621 taa 16619
57 *
dp1ORF131
43846 atgctcaaccggctgagaagaaacttggtggcagaaagatgctactggtttctggtacgctcgagcaaacggaacttatccaa
1 M L N R L R R N L A G R K M L L V S G T L E Q T E L I Q
43930 aagatgagttcgagtatatcgaagaaaacagtccttggttctactttgacgaccaaggctacatgctcgctgagaaatggttga
44013
29 K M S S S I S K K T S L G S T L T T K A T C S L R N G *
dp1ORF132
15304 gtgactggaaggctcatctaatacacatagcctcaagacatttcggttggttttcaggaaaacattcgactagattgtcaattgat
1 V T G R S S N T H S L K T F R W L S G K H S T R L S M Y
15220 cccacaaaggcttcaagggttttcgagttcttcgctggtcctttacagcaagaaggaagtttattcgacctcttgacgttga
15137
29 P T K A S R F S S S S P W S F T A R R K F I R P L A R *
dp1ORF133
8061 atgactttctcattcatgacaagtttttcgagttttcgttggttgctcaggaatagttttcccgccggtcaaaatgtatagattga
1 M T S S F M T S F R V S A C L S G I V F P A A K M Y R L
7977 tcgtatttttcttctgatagcagaactggaatccatttgattcccaccatttcgcccctatctcgccggaataa 7900
29 S Y F S F L I A E L E S I C I P T I S A L S A K *
dp1ORF134
498 atgacttcaatgtacttaggttccatcataatcaagtcattcaaaataatgttcatgcaatcttcgtggaagtccacgtgg
1 M T S M Y L G S I N S Y K S F K I M F M Q S S W K S P W
414 ttacggaaactgaataagtaatttcaatgatttagattcaaccatcttttcggttggaatgtaa 349

388

29 L R K L N K Y N F N D L D S T I F S F G M *
dp1ORF135
780 atgaagcagaacttgaaaatgctgctaattgttgcaatgttctacggagtcaggttcaccattcttgaaattgactcgaaaatctt
1 M K Q N L K M L L M L Q C S T E S S S P F L K L T R K S
864 actcaagctctagctcttcttattacaaggaaaaggcgaatttcacatggaaaatcttacgctgaaatcctag 938
29 T Q A L A L P Y Y K E K A K F H M E N L T L K S *
dp1ORF136
55252 gtgaagaaatcttcaataaccttattcgcttcttgacagatacattcatctgctcagcgattgagttagccccgcggccgtac
1 V K K S S I T L F A S L T D T F I C S A I E L A P R P Y
55168 ataagacctaaaagaacggacttgacagaatttcttcgaagtttcttccctgtagtgcgttcgctcggttag 55094
29 I R P K R T D L T E F L R S F P S L L V V P S G *
dp1ORF137
37146 atgcttcgaactgtttgttagcaccgtcaggaggacaaactagtcgaacccattcacctgcgtctttgataatatctagcgcg
1 M L R T C L L A P S G G Q T S R T H S P A S L I I S S A
37062 acagcgctacagaagaagcaacgtgtttcaacttctcaggcaagccttctgctagttcataccataatgcgtag 36988
29 T A P T E E A T C F N F L G K P S A S S Y H N A *
dp1ORF138
30662 atgactatatcgaagaacaatgtagtcacccggcctatctgtatcttgcgtcgaattcaactcctggaagcataggagcagg
1 M T I S K N N V V I R P I C I L L V K F N S W K H R S R
30578 cgagagctgaaatgtaggaagaatttcttcaattcttccatcattgtcgttcggttagtcattgttcactcctag 30504
29 R E L K C R K N F L Q S V H H C R S F S H V H S *
dp1ORF139
12092 atgatactaaatcactcaactgtttgacctctgataaattcggttcacgcagacacgcgcatttgagcccttttttagatacc
1 M I L N H S T C L T L L I N S F T Q T R A F E P F L D T
12008 tttcgcaaacacctagatgcttccctcactaaaaggctcatgggctcgaagttcttcgaaagacattttcatatag 11934
29 F R K H L D A S L T K R S W A S S S S K D I S T *
dp1ORF140
20562 atgttttcgatatttctcgcgcctaagacttcagcttggtcattgttcactaccattaggtattcattagtaagtgccttagca
1 M F S I F P A P K T S A W S L F T T I R Y S L V S A L A
20646 aagtttgaaaatttcatcttatttcccttattgttttctttatactattattatacaataatgattga 20717
29 K F E N F I L F S L Y L P F F I L L L Y N N D *
dp1ORF141
42922 gtgctaagagttgttagagatatctctaaaacgctcttggttatttcgatttccattcgaataacttatttagtaggacagta
1 V L R V V E I S S K T L L A L F D F H S N N L F S R T V
42838 agcactcgcgtgcacgctgtaataatcgctcgtaagactgctgtgctgtttagccacattggcatagattga 42767
29 S T P L H A V I I V V K T A V S F S H I G I D *
dp1ORF142
31898 gtgactgtcgaagtttctccaacagttctgtcactttacctaagcgctatttagggatttcccggttagcgattaggttcag
1 V T V E V S P N S S V T L P K S V L G I F P L A I R F M
31814 acacctcgtcgaattttaaactggataggttcactaccttttgaaaatcctggaagtgcgatgattga 31743
29 T P A A R I L T W I G S L P F E N P G S A M I *
dp1ORF143
7565 atgaagtttggttgacgcttttaactccagacggttaattttttcaaggcttgaaattggatacatataatcttttcatgc
1 M K F G L T L L T P D R L I F S R L E I G Y H I I F S C
7481 ttttgaaaatacactaaaattccggcgagaataaatttgcatccatctgcgcgtgatagctggaaccattga 7410
29 F W K Y T K I P A R I N L H P S A R D S W N H *
dp1ORF144
36517 gtgcaaatcaagcgactaacttatttagatacattaaacaggcgacattcttcaagattcctaattggaattcaacaattacca
1 V Q I K R L T Y L D T L N E A H S S R F L M E I Q Q L P
36601 ttgaataccgagcgatgacgcagcagcttggaactcttactcttcccgctcaagttgaactgtttctaa 36669
29 L N T E P M T Q Q L G P L L F P L K L N C F *
dp1ORF145
42067 atggaacacgctggagacctaaacaagtggaaagaggttctatttaagcaagacttcgaacagaataattggcagaaactgttc
1 M E T A G D L T S G K R F Y L S K T S N R I I G R N L F
42151 ttcaaagtgggtggaaccatcactcaacctatggcgacgcattctattcgaaaactcttgacggcatag 42219
29 F K V G G T I T Q P M A T H S I R K L L T A *
dp1ORF146
51484 atgacaaaactgcatgattgcatcacctttccagtacggaacctcaagggcgaaacagttattcttcaacgctcgaagtgttcgtt
1 M T N C M I A S P F Q Y G T S R A K Q Y S S T V E V F V
51568 ctaagtttaccagtagcgtgaagatgacctaaaacggaatttctttagggccaatagagcttgtag 51636
29 L S F T S T V K M T L K R N F F M A N M S L *
dp1ORF147
55207 atgtatctgtcaagaagcgaataaaggattttagaagatttcttcaccgagttccctaagtgccagactatatcatattcgttc
1 M Y L S K K R I R L L K I S S P S S L K W Q T I S Y S F
55291 aacagcaggcgaggttggtgatatgttcaaacagctaccggtcgaagaagaggttccctgatatga 55359
29 N S R R R T W D M F K Q L P V E E E G F L I *
dp1ORF148
28636 gtgtttcggttcaagaccattcgagtagggcgaaacacctgtacgattttcgatgtcatccattgctgctaaaatgtcagcgata
1 V F R P K T I R V G R T P V R F S M S S I A A K M -S A I
28552 gggctactttcagctgggttagtccatttcttagtgactgcattgttgcttagcatccatggttag 28484
29 G S L S A G L V H F L V T A Y C C L A S M L *
dp1ORF149
26474 atgccattgaacttttcgagcataaggattaaccttgccccattgtctcactccagctgtggcggaatggcctaaggtagttcg
1 M P L N F S S I R I N L A P L S H S S C G G M A N G S S
26390 agcaagtcaagggcattgtattcgagattttgatatttatgagcagcaggtttccctag 26331

389

29 S K S K G I V F E I L I F M S S R F P *
dp1ORF150
15185 gtggctcctttacagcaagaaggaagtttattcgacctcttgacggtgatagctcttcgcaagttcgacgattcggttggttcat
1 V V L Y S K K E V Y S T S C T L I V F A K F D D S F V H
15101 ttgctttcgctgattgttcgatgaataggtcctcgtatttaattggttcacaagttcgctcgacgtag 15033
29 L L S L I V H A I G S S Y L I V S Q V A S T *
dp1ORF151
28027 atgattatatcaacgcagggagattgctagctacattcaagcacttccttcaaagcgtcttcaataccttggaccaactcttt
1 M I I S T Q G R L L A T F K H F L Q T L F N T L D Q L F
28111 tcctaatgctcaacaaacagggacagacatttcattggtcgaaggtgcaataatttgccagtaa 28176
29 S L M L N K Q G Q T F H G S R V Q I I C Q *
dp1ORF152
42235 atgtgcataaaggacttatcgacaaaagggtactattgcagttacttctgaaggatttagaccgaaagtttcaatgtatcttc
1 M C I K D L S T K R L L L Q Y F L K D L D R K F Q C I F
42319 aggtcttcaataactcatatggaatgccattctatgtatatacactgacggaagacttgggtga 42384
29 R L S I T H M E M P F Y V Y T L T E D L W *
dp1ORF153
22307 atgggtggacaagggtcaccttttcgaactttcgatctcgatagcagacggttccattcggttcaggaataacagttatcgat
1 M V D K G L T F S N F R Y R H S R R F H S F R K N S I D
22391 ggctctttcattttccctttgggcatgacggaattcaacggacaaaactttgccatctggtgta 22456
29 G S F I F P L G H D G I Q R T K L C H L W *
dp1ORF154
18446 gtgacaataggctttaagaactgcaaaaaaacctggggcgtctgcacgcgcaacctggagctccttaacagtcattccaaggtg
1 V T I G F K N C K K T W G V C T R N L E L L N S H P R L
18530 aggtttcttacaacaatcctaattccttcaaatagctcttgcgggtcaatagtcctaa 18592
29 R F L T N N P N S F K I A L V R V N S A *
dp1ORF155
13512 atgaatcagaccctgagcaacttacaatgggacatgggtgcaaaatctaatttccttcttcaacgtttcattcaactcacgacag
1 M N T T L S N L Q W D M V Q N L I S F F N V S F N S R Q
13596 ttgaagctcaagcaattttctggcatatgggagcctatgatattagtccttatgcaaatgga 13658
29 L K L K Q F S G I W E P M I L V L M Q I *
dp1ORF156
18777 atgctagtatctccattttctgttggctctgttttttagctctgttcagttcagctgtcttctcgcatgcaatagtttcgagaat
1 M L V S P F L L V L L F S S V Q F S C F S R C N S F E N
18861 atgctgttcataggtcacaatattccgccaagatttgccagttatggtggcgtcaattaa 18923
29 M P V H R L T I F R Q R F A S Y G G V N *
dp1ORF157
13281 gtgcttgcgtgacttgagaagaattggtatcatttttcgagccaatccataaggttctcgataccgtcacgattgattgtttct
1 V L A G L E K K L V S F S S Q S I R F S I P S R L I V S
13197 gttactgctttcttgaagcgttttttaagtctgtcatattagaccctttcattttctataa 13135
29 V T A F L K R F L K S V I L D P F H F L *
dp1ORF158
40727 gtgaacgcgcttattaggggtcaaacgaagcccaacggacattgtctttgtcccgctcactattgtgaggaacagtcacttctcc
1 V N A V I R V K R S P N G H C L C P V T I V R N S H F S
40643 acttcgagcgttacctcttcgcccagctgtcgtatgctgggtgactgctatgaacacttga 40581
29 T C E R Y L F A G R V V V W V T A M N T *
dp1ORF159
30371 atgatttggctcgcgttacccaagcagcttctcctttgagttctgtcgcagcattccctgtacggctgtccaaatagcatgc
1 M I W S A L T Q A A S P L S F C R A F P V R S V Q I A C
30287 gtctttgcgtattcttccatcttagtagcagcagacttcgcagactgttatgacagcagattga 30225
29 V F A Y S S I L V A A T S Q T V M T A T *
dp1ORF160
41324 atgggttacagacacgcgaggaaaacaatcgaacgtccaagcgtatctatcaatgttatagaatactatggaccgtctatcaa
1 M G Y R H A R K T I E R P R R I Y Q C Y R I L W T V Y Q
41408 ttctccggttcaacgtactcgtcaaatcctgcaattatccaagctcttcgaaatgctaa 41467
29 F L R S T Y S S K S C N Y P S S S K C *
dp1ORF161
52175 atgcaaaaagggttaaatgcttatctcgacatgacattgaaagcattgcattcgagactatttcaaatgtttggcaacggttca
1 M Q K G L N A Y L D M T L K A L H S R L F Q N V W Q R S
52259 aatcaaaccaagggttcaagttttcaacttaccttacaagactcttcaagaatagaatag 52318
29 N Q T K G P S F Q L T L Q D S S R I E *
dp1ORF162
13020 atgacagaagttgcggttaaatagcccgcaaaagggtgagagtagttatgggtcgggaatattgaattttctcgaatatttaaaagg
1 M T E V A V N S P Q K V R V V M V G N I E F L E Y L K R
13104 aagtcaggaacagaaacttccatcagttatattatagaataatgaaagggttcaatatga 13163
29 K Y G T E T S I S Y I I E N E R G L I *
dp1ORF163
40224 gtgaccgaattttctatgtttctccgacgggaatgaagttatgtaccttacgcaagggttcattcacatcgataacgggattcttta
1 V T E F L C S P Q G M K L C T L R K G S F T S I T G S L
40308 cccaatccattcaagtcggccgatttagaacggaacaataactcgtttaatccagacatga 40367
29 P N P F K S A D L E R N N T R L I Q T *
dp1ORF164
6696 atgtactcttggagaacttcgtgcctaaatgttcagcttcgcccattgcaattaggttagaatctcggttatctataatagac
1 M Y S W R T S C L N V P A S P I A I R L E S A L S I I D
6612 tcacgattcttccgaaatacatttttcgaatacaccaccaaccccgctgggttataa 6553
29 S P I L S K Y I F R I H P P T P L G L *

390

dp1ORF165
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1 M S E S W S I P T T D G L Y L D I M L S K I A G V R F F
50420 cctccaatcataaagggcgtgactaccacaaggaattttcagcctcagtcattgcttga 50361
29 P P I I K G V T T T R E F S A S V I A *
dp1ORF166
23519 gtggctcatgctctttaatgactctatcttctcccgcttggctcgctttactgtcccagctgtaagcatagattcatcaatgtc
1 V V M L F N D S I F S R L A R F T V P A V S I V F I N V
23435 gtgcgtgttgcagggtcgagtgtaaatctattctcagccaagagttcagcgtgaaatga 23376
29 V R V A R V E C K S I L S Q E F S V K *
dp1ORF167
1008 atgcttattcggttggagcttcttactcgtctatattgggtctcacgcagacgatgcggctggaggtgcttaccctgattgcactc
1 M L I R L E L L T S Y M V L T Q T M R L E V L T L I A L
1092 ctgagttctataattcaatgtcaaatgcaatggaatggaactggaggcaaggtaa 1148
29 L S S I I Q C Q M Q W N M E L E A R *
dp1ORF168
54345 atgagactttttccaggttatattcttcacattgttcagttcctcgagtcaggtattgttcttgaaattcatagagttcgaaag
1 M R L F P G Y I L H I V Q F L E S S I V L E I H R V R K
54261 ttgtgcaagggtcatagggccgacatcatataggcaacatcaggaggaaattaaactaa 54205
29 F A K G H R P H T Y R Q H Q E E L N *
dp1ORF169
45954 atgaacacagcatcgcaagagtttcaatgttagtgataaggaagaattcgtcggtggccaccaagcaagcttctgcccgttta
1 M N T A S R R V S M L V I R K N S S W P P S K S S A R L
45870 gaaactccgtcaatcactaatttcccatctttagtgactcgacttctctaaaatga 45814
29 E T P S I T N F P S L V T R L P K I *
dp1ORF170
27600 atgatgattgttcttctgtgctcctcgcttgggtgagcagcagcaagttgcttaccaaaagagccgatttcacgaggttcgggaa
1 M M I V L V L L P F V E Q Q Q V A Y Q K S R F H E V R E
27516 caccaccaccgacacgactggatttccataatttccagtcgccggtggcgacttag 27460
29 H H H R H D L D F L N F Q S R L A T *
dp1ORF171
47678 atgtcattttcttctcatgtactcttttagagcatcacgaagacttttgacttgtttctccatgtcgcctttggtagcattta
1 M S F S F M Y S F R A S R R L L T C F S M S P L V A F N
47594 tcaccggcttcttcaattgcagcgatgaactgttttctcatcttcaatttcatctaa 47538
29 S P A S S I A A M N C F S S S N F I *
dp1ORF172
10462 atgtttcgaacattttctacccattattagaagcagcatcaatttcaataggagagccaagtcctttgttcacatccttcg
1 M F R T F S T P L L E A A S I S I G E P S P L F T S F A
10378 aaaattcgagcagtagtggtttaccagttccagcgccaccacagaatagatag 10325
29 K I R A V V V L P V P A P P Q N R *
dp1ORF173
32160 atgacattagacatttcttctcgtctgtacgaaagggttccagctttagtcaacttcaccgtacattgcactgaagattgtcataag
1 M T L D I S F V C T K G F S L S H F T V H C T E D C H K
32076 ttgctcatctgtcatatactcgccgacttcagcgraagtaggctctaccattga 32023
29 L L I C H I L A D F S V S R L Y H *
dp1ORF174
29766 atgtcccatcagcccttttccattaagattgtcgaaccagcgttcgacttttcatcagtttcaagctgttcttgcattatattggt
1 M S H Q P F S L R L S N Q R S T F H Q F Q A V L A Y I G
29682 cataatagaattgcgccatttgtttccagtagtctgcgtcaccttttagactga 29629
29 H N R I A P F V S S S L R H L L D *
dp1ORF175
15648 atgcgcgtgatgtcatggcagataggcgaggataaagagtgctgaatagaacgccgagagcttacgagagcgccaaatacaag
1 M R V M S W Q I G E D K E C R I E R R R A Y E S A K Y K
15564 ggcgacggtactacggtggtcctcttgcattacgtgaacaaataaaccattga 15511
29 G D G T T V V L L L T C N Q I N H *
dp1ORF176
43031 gtgataaagacggtaacgttgaatttttctagttccgtcttgaatgacgtcattttgggtgattgattgctactgtcgtttggtc
1 V I K T V T L N F S S S V L N D V I L V I D C Y C R L V
42947 aatcccgctcgacctgctgtttaagagtgctaagagttgttagagatatcctctaa 42894
29 N P V D L L F K S A K S C R D I L *
dp1ORF177
19937 atgaacctaaacagttcgagacttctcaagctgttgggaaagaagcaggtcgaatattttgggtgggaacgtgaacttggtcata
1 M N L N S S R L L K L G K K Q V E Y F G G N V N L V I
19853 ttctcgcgactaatttttaggtgcttttgtattaatcagcgtgatgacgcttga 19800
29 F S R L I L G A F V L I S V I C A *
dp1ORF178
11924 atgacaactgtcgaccaattttaaagacagttgaggaaaagtttaggtcaatttttcttcatcagtttctttaaatttgagc
1 M T T V D Q F K R Q L R K S L G S I F P S S V - S L N - L S
11840 caattagtaacctttagegaattgctagcacttgcctcccatattaaagtcataa 11787
29 Q L V T F S E L L A L A S H I K S *
dp1ORF179
56058 atgggtagggttattccttaccctcggtgatttgcctttatgcgaaacacctaccacaatcgcttgcgtggcttcaggagttgcatt
1 M G R V I P Y L V D L L Y A K P T T I A C R G F R S C I
56142 ttggataagtcaaaaagcaagtgcttttatattcgacaagctctcgaataa 56192

391

29 L D K S K S K C L Y I R Q A L E *
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41176 atgttcgacatgatttggaggaagtgttccctgttaaaatctgtcgaaccgctgaagtcgtgtctactaaagaatgcccgaa
1 M F D M I W R K L F P V K I C R T A E V V S T K E M P E
41092 aaagtaggacgtactgaatcggggatgttgaacctccatccgtttgaatag 41042
29 K V G R T E S G M L N L H P F E *
dp1ORF181
13126 atggaagtttctgttcgacttctcttttaaatattcgagaaattcaatattcccgaccataactactctcaccttttgcggg
1 M E V S V P Y F L F K Y S R N S I F P T I T T L T F C G
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29 L F T A T S V I G C P P L L I L *
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45369 gtgcttgcccatgtttcaataaatagggttcgacctcgcttagctttcgaacgtgctataacgatttcaatcatagcgaagaaa
1 V L A H V S I N R V R P R L A F E R A I T I S I I A K K
45285 ggtgagaagcttcaatcaattccattgcggtgtcaatatcttcttctctga 45235
29 G E K L Q S I P L R C Q Y L L P *
dp1ORF183
13896 gtgattccagcttttggttttcttcagcctcttcaacttttcttcttaggcgcaggtttcttacgagttgaactcttaggt
1 V I P A F G F S S A S T T F S S L G A G F L R V E L L G
13812 ttttcttcaactacttcttcaacctcagcctcttgttcaactggacctga 13762
29 F S S T T S S T S A S C S T G P *
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53330 gtgaacttgcgctcaaccacgtcaaacatttggctctcgtaggtctaaaattagagttccaagaagttcgctcttttctgga
1 V N L P S T T S N I W S S S R S K I R V P R S S L F S G
53246 aaatcttcaagagtagcactgtcttccgacgtctggaaggaattcataa 53196
29 K S S R V A L S S G R S G R N S *

dp1ORF185
22522 atgaaattcgagatgttcgaaatgaaatctactattattagacacttttagaaatggcgaagaattgtcaactacttctata
1 M K F E M F E M K I Y L L L D T L E M A K K L S T T S I
22606 tatttggaggaaaagatgagtcgagtcgaagccttatacagggggtga 22653
29 Y L E E K M S R V K T L Y R G *
dp1ORF186
21272 atgctcgaaaaactcaaccggttcgaaaacctcaatccttcgaaaagtcgaaccattcgaaaagttcaaaagttcgaaaaactc
1 M L E K L N R F E N L N P S K S R T I R K V Q K F E K L
21356 aaccttcgagagtaggaattaaggacataccagttcaaccttttag 21403
29 N H S R V G I K D I P V Q P F *
dp1ORF187
34415 atggcttctgttcaatctcttctactatcattcaagcagctgttcaaatatcactgctttattcaatggtcttgttcaggcac
1 M V L F N L F L L S F K Q L F K L S L L Y S M V L F R H
34499 ttcctacgcttattcaagcaggtcttcaaatttgtcagctctcataa 34546
29 F L R L F K Q V F K F C Q L S *
dp1ORF188
35609 atgttcgtaaaagcagccggttcgcctcgagtggttcaatacaggaagtgacaaccttaaccaacctcagtcacaatcta
1 M F V K Q P V R L E W T C S I Q E V T T L T N L S H N L
35693 aaaacaatcaaggcgagcaaacggttgcacattggaacaatcgtag 35740
29 K T I K A S K P L S T L E Q S *
dp1ORF189
42587 atgcaaacgcagtatcaaccgtctctgaaactcttcatgaccagacttgatgctgcaaacgctcgagaacttcgagctgacg
1 M Q T Q Y Q P S L K L F M T Q T C M L R T V E N F E L T
42671 agcaaaaacttcgcgaactcgttacgcaatcgaagatgaaattctag 42718
29 S K N F A K L V T Q S K M K F *
dp1ORF190
39786 atgtattcactcaaaagttgttcagtggtcgaatcatattaaaaatcgaaacttggttaatatcttactccttttagtgaagcag
1 M Y S L K V V Q C G S I I L K S N L V I S L L L L V K Q
39870 aggaagaccttaaatatcgaattgactcaaaagccgatcaaaagctaa 39917
29 R K T L N I E L T Q K P I K S *
dp1ORF191
40996 atgtccattgttcggaacttgatttaggtaagtaccttgctaagtcagtgacggcgtaaaaggatacgctagtagtatgggttc
1 M S I V P E L D L G K Y L A K S S D G V K D T L V V W F
40912 ttacctaaatctatccagtcgctaccgaaaactcgggtaccaaacttga 40865
29 L P K S I Q S L P K T R Y Q T *
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1 M V D V E C F F E M K F R V F S I P Y G M F S E C P N K
2836 acggaatggagtatcttgcacccgtcacgttctgcgtcctcgctaa 2789
29 T E W S I L Q P V T F C V L A *
dp1ORF193
42456 atgatttcagctcaaatcaaatcgaagaatgagacattgttcaaatcaaccaagaattatctacattcgatttcacacaaagtc
1 M I S A Q I K Y E M R H C L N L T K N Y L H S L S P Q V
42372 ttccgtcagtgatatatcatagaatggcatttccatgatgatttga 42325
29 F R Q C I Y I E W H P H M S Y *
dp1ORF194
40284 atgaaccttgcgtaaggatcataaacttcattccctcgaggagaacatagaatttcgggtcacttgataccttaatggtagacta
1 M N P C V R Y I T S F P A E N I E I R S L D T L M V E L

392

40200 cgcgcgttcttaccgataaattagaccttcattagaagagctcatgtaa 40153
29 P S F L P I I R P S L E E L M *
dp1ORF195
42584 atgttcacaatcggtgttttgacaagtttcttttcagctccttgccaatagtgaaactctgccacaatttggcgcgattttgta
1 M F T I V V L T S F F S A P C P I V N S A T I W R D F V
42500 aggttcaacatagttctcacctcctttctaaaaatattataacatga 42453
29 R F N I V L T S F L K N I I T *
dp1ORF196
11273 atggttagatttaacaagtcctctgccaatcatgtcactcctccttgctcatcaaaagaagtttggtttcaattatcggttttagc
1 M V D L T S P C P I M S L L L A H Q K K F G F N Y R F S
11189 attaggctcccatatttaacaactccagcaagttcattcatttctcttag 11142
29 I R L P F N N S S K F I H F F *
dp1ORF197
7484 atgaaaagatttatatggatccaatttcaagccttgaaaaaattaaacgggtctggagttaaaagcgtcaacccaaacttcatcg
1 M K R L Y G I Q F Q A L K K L N G L E L K A S T Q T S S
7568 atgcagggtatgaagtttcttacaagaagcgtcgaaactagattga 7612
29 M Q G M K F L T R S V E L D *

dp1ORF198
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1 M P L N K L T S S F I Q C L S S P I Q L T L E T L P A C
24203 tttctgttgacattgtttatcaggacgagcgtacaaaaggaaatga 24247
29 F L L T L F I R T S V Q K E *
dp1ORF199
15742 gtggctcctgaattagcgtgtacttttctcccaactgccttagcaactgccttctctgttttagcactagctctgcgcgtggga
1 V A P E L G C T F P P N C L A T A F S C L A L A L R V G
15658 attggtttgtatgcgcgtgatgtcatggcagatagggcaggataa 15614
29 I G L Y A R D V M A D R R G *
dp1ORF200
47843 atgacaggcttgattcgataagccctgaaagtttttcacacatttcttccgtctcggcttcgtcaactaattttcgataatt
1 M T G L Y S I S P E S F S H I S S V S A S S T N F S I I
47759 tctttcaagcgttcttccatagttgagcgtctgtcgtgtag 47715
29 S F K R S S S I V E R S V V *
dp1ORF201
38569 atgggcttcacaagttccttcttatacaaaaggtcaatatcttggactcgaactatttggacctataccgattcaactaccga
1 M G F T S S F P N Q R S I S L D S N Y L D L Y R F N Y R
38653 aacgggctatcaaaaaacctacattccaaaagacgggaatga 38694
29 N G L S K N L H S K R R E *
dp1ORF202
44483 gtggggcggtttattttttataaaaaattttttacaaaatgcttgacaacattcactcattatcgatataatacaattataaaaaata
1 V G R L F F I K I F Y K M L D N I H S L S Y N T I I K I
44567 aataaagccgaaaggcgaggacattatgtcaaaaattaa 44608
29 N K A E R R G G H Y V K N *
dp1ORF203
22781 gtgattaggattggccgggttacaagagaaccacattttcgaacctgttacggaacagcgcctgtcgttgggtgacaaacga
1 V I R I G R V T R E P H F R T C Y G T A P C R L V D K R
22697 ttcaggcatcagtgccacctcatcacagaagatacctgctaa 22656
29 F R H Q C H L I T E D T C *
dp1ORF204
1471 atgaccacgggttcgagtcagggtggttggtagcttttactcagtcagcaagaaaatcgaggtacattcattgacagacttgacc
1 M T T V R V K G W L L T F T I T S R K S Q V H S L T D L T
1555 acgctgttcttcttcaagggaatgaaccaatcgcttag 1593
29 T L F F F K G M N Q S L *
dp1ORF205
8524 gtgacactgatgaatgggttctcagtttggtatgctactcgtgacgcagatatcttctacgaccaaaagaattgccaatttagaa
1 V T L M N G S Q F G M L L V T Q I S S T T K E L P N L E
8608 ttcaggaaaagcaacctgctatcaagttcaatttcgtag 8646
29 F R K S N L L S S S I S *
dp1ORF206
19855 atgaccaagttcacgttcccaacaaaatattcgacctgcttcttcccaacagcttgagaagtcctgaactgtttaggttcac
1 M T K F T F P P K Y S T C F F P N S L R S L E L F R F I
19939 aaattgttcaacttgagcaagtgcgatattattctttag 19977
29 K L F N L S K C D I I L *
dp1ORF207
27502 gtgtcggtggtggtgttccgaacctcgtgaaatcggtccttttggttaagcaactgctgctgctcaacaaacggcaggagcac
1 V S V V V F P N L V K S A L L V S N L L L L N K R Q E H
27586 aagaacaatcatcattctttaataataggaggaactaa 27624
29 K N N H H S L N N R R N *
dp1ORF208
47279 atgtttggtatgaagcaaaagacttcgctgaagaaaataacattcacttcccggttgggttcttctgaacctagaacagacctg
1 M F G M K Q K T S L K K I T F T S R L F F L N L E Q T L
47363 accatcggttctcgtggttggtatgacgaaggcgtga 47401
29 T I V V L D S G M T K A *
dp1ORF209
29784 atgttaagaatcaagttcgtagaccattgaaacctcctactaaaatcaaggtacttcgaaactcttgggtcagtgatggat

393

1 M L R I K F V E P L K P L L L K S R Y F E T L G S V M D
29868 atggaggagaaagaaaaggataaaagcgaatgaagtcgtag 29906
29 M E E R K R I K R M K S *
dp1ORF210
53077 atgtttcaacttttccccgtatcatggttgtaaagttgaagaaatagttttcaatacagagggaatccggttttggcataatggac
1 M F Q L F P Y H G C K V E E I V F Q Y E G I R F G I M D
52993 aattatcaggatggactgtttccccgtcttcgccaatag 52955
29 N Y Q D G L F P R L R Q *

dp1ORF211
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20875 ttacttattaagtccttttctatattagattgtttataa 20837
29 L L I K S F S I L D C L *
dp1ORF212
52983 atggactgtttccccgtcttcgccaatagcattgcaattgatatagcgtcgacgaccgtcaacgtctgcttcggtgactacgaa
1 M D C F P V F A N S I A I D I A S T T V N V C F V D Y E
52899 ataaccatgtcttcgccttcgggtcatcatacaatag 52851
29 I I H V F A F R V I I Q *
dp1ORF213
30291 atgcgtctttgcgtattcttccattcttagtagcagcagacttcgagactgttatgacagcgacttgaaactgttttcgataccg
1 M R L C V F P H L S S S D F A D C Y D S D L K L V S I P
30207 ttacaggttactaacaattcttccaggttccatactaa 30169
29 F T V T N K F F R L P Y *
dp1ORF214
24273 atgatgccaaagtgtttttcagtgctcattccttttgtagcgtcgtcctgataaacaatgtcaacagaaagcaagctggaagg
1 M M P K L F F S A H S F C T L V L I N N V N R K Q A G R
24189 gtttctagggtcaactgtataggtgaactgaggtcattga 24151
29 V S R V N C I G E L R H *
dp1ORF215
35822 atgttaccacacccgtgatagagtcttcttacttctattatacaatcctctcgacagtttgtaacgtcgtcattgtttcgaact
1 M L P N P D R V S L L L L Y N P L D S L S T S S L F R T
35738 acgattgttccaatgttgacaacgggttgctcgcttga 35700
29 T I V P M L T T V C S P *
dp1ORF216
32849 atggcctcggagctcggggccacatctcctccagatacggcgagccagggtcaagtacccctggcatagcgtccatgatttcattt
1 M A S E L A A T S P P D T A A R S S T P G I A S M I S F
32765 acctggaacccgggtgaagctagattttccataccttga 32727
29 T W K P A E A R F S I P *
dp1ORF217
23443 atgaatactatgcttacagctgggacagtaaagcgagccaaacgggagaagatagagtcattaaagagcatgaccactgcatgg
1 M N T M L T A G T V K R A K R E K I E S L K S M T T A W
23527 ataggaacagatatgctgtctcactgacgtctaa 23562
29 I G T D M P V S L T L *
dp1ORF218
22029 atggaatgcttccggaagaggttcgatataactacaattgagcgcgagaaaattacattgctccgggcaaaaatggcgacc
1 M E C F R K R F D I D Y K L S A R K L H C S G P K W A T
22113 aggaattgaaggcgaggttaagataacttcgtag 22148
29 R K L K A R L K I T S *
dp1ORF219
51388 atgattttatgctcgactttttcagttctccatttcttcgaaaacgttccagggtgacgccttgcttaactacttcgctagat
1 M I L C S T F S V L P F L R N A S G L T P C L T T S L D
51304 gttccaaaattccttttcagccactgggttccatag 51269
29 V P K F L F S H W F P *
dp1ORF220
6334 gtgaagttttcttcgggtgacggttgatacaatttccttcaagagtaagctgttaaggtggcaagtgattctttcttcgaaact
1 V K F S S V T V D T I S F K S K L L R W Q V N S F F E T
6250 ttcttgccagcagatgctgacatgatgtcttcataa 6215
29 F L P A D A Y M M S S *
dp1ORF221
43507 atgactgctcaagttctatgtactatgctctccgctcagccggagcttcaagtgtggtgggagtcataactgagtcacatgc
1 M T A Q V L C T M L S A Q P E L Q V L D G Q S I L S T C
43591 acgcatggcttattgaaaacgggttatgaactaa 43623
29 T H G L L K T V M N *
dp1ORF222
13212 gtgacggtatcgagaaccttatggattggctcgaaaatgataccaatttcttctcaagtcagcaagcactcgataccatggaa
1 V T V S R T L W I G S K M I P I S S Q V Q Q A L D T M E
13296 gctatgaaggtggacttgcgagcactcattaa 13328
29 A M K V D L S S T H *
dp1ORF223
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1 M W W Y L L D M F E M S T T S T V K S L T F T T R K M S
14139 acgagcctgacgatgacagcgacattcttctgtag 14171
29 T S L T M T A T F L *

dp1ORF224
13621 atgccagaaaattgcttgagcttcaactggcgtgagttgaatgaaacgttgaaagaaggaaattagattttgaccatgtcccat
1 M P E N C L S F N W R E L N E T L K K E I R F C T M S H
13537 tgaagttgctcagggctgattcatatgctaa 13505
29 C K L L R V V F I C *

dp1ORF225
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1 V S N G C D V F H R L C H V A S F C V R I S C C S S K Y
32907 gtcagccacgtgacccgctggtttgcctctaa 32875
29 V S H V T R L V C L *

dp1ORF226
25191 gtggctgctacattagtttgaacttcagtgagcgcaagtgtgcttagcagaagttcatcgctaggaattggatagtggtgttc
1 V A A Y I S L N F S E R K L L S R K F I A R N W I V V F
25107 gatagtcattgtcgtgaagtgtttgataacttga 25075
29 D S H C R K C L I T *

dp1ORF227
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1 M T Q L D G S A Y D V S R I H K G R R L L H Y R Y Q S R
23031 ctgctacgaataaacggtcgaattctatatga 22999
29 L L R I N G R I L Y *

dp1ORF228
10450 atgttcgaaacattattgaagattctagatacaagtctatggacagcgagttcaaagtttacatcattgacgaggttcattatgc
1 M F E T L L K I L D T S L W T A S S K F T S L T R F I C
10534 tttcaaccggagcatttaaatgcgctgttga 10563
29 F Q P E H L M R C *

dp1ORF229
27634 atgtgcgagtttaagaaaactgatttttaatacaaacctcgaagcattgtcgcaattcctgaccactacgttgcgttggctgctc
1 M C E L R K L I L I K P L E A L S Q F L T T T L L W L L
27718 aaattccagctaccgcagcaactcaagtag 27747
29 K F Q L P Q Q L K *

dp1ORF230
50723 gtgacgaaaaatccggcactactgaactatctgtcgttaaaaaccgatatggcgaagaccgaaaaatcatcgaatatatgtggg
1 V T K N P A Y L N Y L S L K T D M A K T E K S S N I C G
50807 acgttgaaactggaacctatactcttatag 50836
29 T L K L E P I L L *

dp1ORF231
31071 atgcgcgtgtcattgcgtttcacatcttcagttccctccgaggtcacggcttcgagttctgctgtttctgccgtatctacgaca
1 M R V S L R F T S S V P S E V T A S S S A V S A V S T T
30987 aagttagctccgcgacttttggcaactga 30958
29 K L A P P T F G N *

dp1ORF232
29385 atgtcaattccattagctcttgcataattcaacgagctcaggaacgggttttagccgcatactcttcgcgcatttgttcaacttcg
1 M S I P L A L A N S T S S G T V L A A Y S S R I C S T S
29301 tcaatttcttcaactgattcaattgtttga 29272
29 S I S S T D S I V *

dp1ORF233
52892 atgtcttcgccttccgggtcatcatacaatagagtgaattgcgctgtcaccgtggtcagcgagtgtaaaaaactcgttatta
1 M S S P S G S S Y N R V T I A L S P W S A S V K N S L L
52808 gaccctgagctaaatgttccgtgattttga 52779
29 D P E L N V P D F *

dp1ORF234
36253 atgcttacgagtagacgactcaactgttcgaaagggtttataagtttcaaccgcgtttggaggcgatagcttacctaaccag
1 M L T S T A T Q L F E R F I S F N P L W E A I A Y L T Q
36337 gaagacctactcgacaatttagagtag 36363
29 E D L L D N L E *

dp1ORF235
32768 atgaaatcatggacgctatgccaggggtacttgacctggctgcgctatctggaggagatgtggccgcgagctccgaggccatgg
1 M K S W T L C Q G Y L T W L P Y L E E M W P R A P R P W
32852 ctagtctcaactcgagcctttggattag 32878
29 L V H F E P L D *

dp1ORF236
37528 atgttcgtcgttttagatttagcaatatatcgaggcttcgtggtgtagtaaacacgaacatcaatgagatattcact
1 M F V A F R F S N I S R L H V A C S K P R N I N E I F T
37444 tccattgtgatagaagcaaacgttaa 37418
29 S I V D R S K R *

dp1ORF237
1678 gtgagagtcaggttaaggaactcttgacatattctcagccgtagttctaaatccaaatagaactcgcttggtgtcaactgcattt
1 V R V Q V R N L D I F S A V V L N P N R T R L V S T A F
1594 gctaaagcgattgggttcattcccttga 1568
29 A K A I G S F P *

dp1ORF238
1301 atgcctttttgcggtcgatacaagttgcgcaagttccacaactttcagcgtcactttcataacatgaacgagtcagaataaag
1 M P F C G R Y K L R K F H N F Q R H F H N M N E S R N K

1217 gaacatctaaatcaattccccatttaa 1191
29 E H L N Q F P I *
dp1ORF239
26521 atgggtgaagtatttcctatcgagaatgtcctttcgaccatcctaataatggaatgtgctaccaaactgtatggtacgaaaactcac
1 M V K Y F L S K N V L S T I L M E C A T K L Y G T K T H
26605 tcgaagaaatcgctgatgagttga 26628
29 S K K S L M S *
dp1ORF240
41893 atgtttggaataagcgtgaaacagagttttacatggcgaagtaacaaatcaggagacaaccctacgggaactcgaggtgaatggg
1 M F G I S V K Q S L H G E V T N T R T T L R E L E V N G
41977 gactatttcaaaatttctggtag 42000
29 D Y F K I S G *
dp1ORF241
47020 gtgtctttccttaatatggagatagttttcattctatttaagcaggatatacgaaaaggttaccaatttttagatttcattaggtt
1 V S F L N M E I V F I L F K Q D I E K V T N F R F H R L
46936 accatctacgatataatctgctaa 46913
29 T I Y D I I C *
dp1ORF242
41338 gtgtctgttaacccatgctcttacggttagcgagccattaaagttcatcatacccaatttgcgcgcttttcggttgatagcttgg
1 V S V T H A L T V A E P L K F I I P N L P P F S L I A W
41254 tttttacctacgagctcagcgtga 41231
29 F L P T S S A *
dp1ORF243
51306 atgttccaaaattccttttcagccactggtttccatagaacctccatcggtttcgacctataacattcgagacgaattcagtta
1 M F Q N S F S A T G F H R T L H R F D L I H S R R I Q L
51222 gtcttgaaagtgtagcgcgaagtga 51199
29 V L K C S R K *
dp1ORF244
27083 gtgaggtacaaaatgttgaccgtcgccgtcaatgaaaatttttagcatcgagttctttcgaaagttttcgaaataatttccttcac
1 V R Y K M L T V A V N E N F S I E F F R S F R N N F L H
26999 ctgtttgatagttggttcatttag 26976
29 L F D S W F I *
dp1ORF245
6278 gtggcaagtgaattctttctcgaaactttcttgccagcagatgcgtacatgatgtcttcataactgctagtagaagttttaat
1 V A S E F F L R N F L A S R C V H D V F I T A S R S F N
6194 tcgaagtcggtctttcaagaataa 6171
29 S K S V F Q E *
dp1ORF246
2831 atggagtattcttgaacccgtcagcttctgcgtcctcgccataatagacaaaaagtccttgaacggctgcctcagttattgtcca
1 M E Y L A T R H V L R P R L I D Q K V F E R L P Q Y C P
2747 aggttacaaatttcacccggttaa 2724
29 R L Q F H P A *
dp1ORF247
29641 gtgacgcagactactggaacaaatggcgcaattctattatgaccaatataagcaagaacagcttgaaactgatgaaagtga
1 V T Q T T G N K W R N S I M T N I S K N S L K L M K S R
29725 acgctggttcgacaattctaa 29745
29 T L V R Q S *
dp1ORF248
53560 gtgcaaagcctcggttctagcaagaagaacgatgctcagttacttgcctcaacggaaaaacaggaagcctgcagttgaggttactt
1 V Q S L V L A R R T M L S Y L L N G K T G S L Q L R L L
53644 acatttcaggaaacgctctaa 53664
29 T F Q E T L *
dp1ORF249
2012 gtggatgcgactatcattgcaactggtgtgactcagcctttacctggaacggtactactgagccggaatatatcacaggcaaaag
1 V D A T I I A T G V T Q P L P G T V L L S R N I S Q A K
2096 aagctgtagtcgaatcttga 2116
29 K L L V E S *
dp1ORF250
23837 atgggcaaacatggaagattgacgaagactcagtcgactataaacctactcgagaaattcgaaactatattcgacaacttatca
1 M G K H G R L T K T Q S T I N L L E K F E T I F D N L S
23921 aaaagcaatcacgctttatga 23941
29 K S N H A L *
dp1ORF251
39205 atggaaataaattagttaccgtctgcgctggttcccggttatcccttgagctccgtcattcccttccatttcgtccatgt
1 M E I I S L T V C A W L P G Y P L S S V I P L P F R P C
39121 ataggctgcagggtcttttga 39101
29 I G C R V F *
dp1ORF252
54771 gtgttgataggtcgaaactaattttgcatattttctatatttcaaaagtgcttttgagatatacgttataaaatgctcgacaa
1 V L Y R S K L I L H I F Y I S K V L L R Y R Y Q N A R Q
54687 tactttcgctgttctcttag 54667
29 Y F R L F L *
dp1ORF253
56255 atggttgcgtctataatagaaccgatgttgtagacaaagcatttgcaatcttcgagtcataatttattcgagagcttgctgaat

396

1 M V A S I I E P M L L D K A F A I F E S N L F E S L S N
56171 ataaagacacttgctttttga 56151
29 I K T L A F *
dp1ORF254
48479 atgaacctttcgcttaggttcaatctttttcgaaacattttcatatttaacaaaactttcagctaaaaatcgacaaagtccaatg
1 M N L S L R F N L F R T F S Y L T K L S A K N R Q S S M
48395 ttcgactcaatgttttaataa 48375
29 F D S M F K *
dp1ORF255
9572 atgcttttggtcttctcgacgaatgactctactacattccctgcagggtttcgagcagtcagggtcaatgatgcaccgttttcgt
1 M L W S S R R M T L L H S L Q G F E Q Y G S M M H R F R
9488 caaggtagtcaccttttctaa 9468
29 Q G S H L F *
dp1ORF256
15289 atgaccttcagtcactaatgcggccgctgaaattggataccactatacatgggttcaccaacttcgagacaaaagcagttgaaa
1 M T F Q S L M R P L K L D T T I H G F T N F E T K Q L K
15373 cacttgaagaaatttttag 15390
29 H L K K F *
dp1ORF257
28216 gtgaacgtgctggatttagcaaaacagctactgagatggcattcttccgtgagtcctatgcgacttggtgaaaaagaccgtcaaa
1 V N V L D L A N K L L R W H S S V S L C D L V K K T V K
28300 acttgcaaatgctattga 28317
29 T C K C Y *
dp1ORF258
44023 atggaaattggtattggttcgaccgtgacggatacatggctacgtcatggaacggattggcgagtcaggtactacttcaatc
1 M E I G I G S T V T D T W L R H G N G L A S H G T T S I
44107 gcgatgggttcaatggtaa 44124
29 A M V Q W *
dp1ORF259
4298 atgactcgactacgaagcataaagacaagtggatggaaagagtattcgaaatttcgaaacagttctaaccagacgttaaga
1 M T R L R S I K T S G W K E Y S K L F E T V L I Q T L R
4382 ctcacgcatttggatga 4399
29 L T H L G *
dp1ORF260
24746 gtgacctacttccctcaatcgccgtactggaggcaagcaagctcaagtcacttccatttcaggaaacttcaacttccctccag
1 V T L L P Q S A V L E A S K L K S L P F Q E T S T S F Q
24830 cggctgaatattatttag 24847
29 R L N I I *
dp1ORF261
288 atgaattcacttccctttgccttaaaacaggacagcctgacttcgcgaatgttttcattagttacattccaaacgaaaagatgg
1 M N S L P F A L K Q D S L T S R M F S L V T F Q T K R W
372 ttgaattcaaatcattga 389
29 L N L N H *
dp1ORF262
9408 atgcctattcaactccaggcggaagatgtggaagcatgcttgcagttcgacttaaaatttagaaaaggtgactaccttgacg
1 M P I Q L Q A E R C G S M L V Q F D L N L E K V T T L T
9492 aaaacggtgcatcattga 9509
29 K T V H H *
dp1ORF263
27052 atgaaaatttttagcatcgagttcttttcgaagtttttcgaaataatttcccttcacctgtttgatagttggttcatctagacctttt
1 M K I L A S S F E V F E I I S F T C L I V G S S R P F
26968 aacaagttctctaattga 26951
29 N K S S N *
dp1ORF264
6139 gtgaatagtacaaggcggtctaatcgcctcaggattttctgctgtagggatagccgcatcatcttcaactcaattgagtcaagc
1 V N S T R R S N T L R I S A V G I A A S S S N S I E S S
6055 tgtgaaacgtcttcataa 6038
29 C E T S S *
dp1ORF265
4801 gtgaataaagtcaagcggtttttgtataaaaagttcatttttttttaaaaaaataagagcgaaaagctcttatctaaaatagtc
1 V N K V K R F C I K S S F F F K K N K S E K L L S K I V
4717 gacgttgacgatttttaa 4700
29 D V D D F *
dp1ORF266
50220 atgcccggttcttccaagcagttgcaagcattttatcaatagtcacgacttaccttgcagggtcgagccattatgacaatcaa
1 M P V L P S S C K H F I N S P R L T L S R S S H Y D N Q
50136 atcctcaccaggaagtaa 50119
29 I L T R K *
dp1ORF267
47367 atggtcaaggtctgttctaggttcaggaagaacaaacgggaagtgaatgttattttcttcagcgaagtcttttgcctcatacca
1 M V K V C S R F R K N K R E V N V I F F S E V F C F I P
47283 aacattaatcgtagatag 47266
29 N I N R R *
dp1ORF268

397

12621 atgtcaatttcggtcttctgtgcttgacaatggattcaactactgatgcgtcaacctttttcaatcgcgacagcttgccaattca
1 M S I S V L C L T M D S T T D A S T F F N R D S L S N S
12537 ttgtcaattcttagagtaa 12520
29 L S I L E *
dp1ORF269
53834 gtgaatagtagtgcagtcacatcagtttctacgtcaatagaacctattccgtcttcaatcattttgtctacatactgctcgagttt
1 V N S I E S I S F Y V N R T Y S V P N H F V Y I L L E F
53750 tgccttcctcagtgattaa 53733
29 C F L S D *
dp1ORF270
50792 atgatttttcggtcttcgcatatcggtttttaacgacagatagttcaagtatgccggatttttcgtcacgcttcatagcgata
1 M I F R S S P Y R F L T T D S S S M P D F S S R F I A I
50708 actctgctagcattttga 50691
29 T L L A F *
dp1ORF271
19739 atgaggctgctttgctttatcttcggtaccgtattgaccgacttctactcgcgaaaccttcctacaagaattcatacctcaaag
1 M R L L C F I P V T V L T D F L L A N L P T R I H T S K
19655 gctttttgtcagccttag 19638
29 A F C Q P *
dp1ORF272
1556 gtggtcaagtcgtcaatgaatgtacctgcgattttcttgacgtgataaaagtcaacaacctcccttgactcgaaccgtggtc
1 V V K S V N E C T C D F L D V I K V N N H P L T R T V V
1472 ataagttccgctgctaa 1455
29 I S S A C *
dp1ORF273
56256 atggatttcattaggactgagtcctcttggaattggaacggttgcatatatagatattccgtcagccgtactaggccaagttct
1 M D F I R T E S S W N W N G C I Y R Y S V S R T R P S S
56340 agttcagtttatcttgagtcgaattgcttcgagatatttgaaaaagtagtcaggaaaattcctgattatcttgagtcgaattgc
29 S S V Y L A V N C F E I F E K V V R K I P D Y L A V N C
56424 ttcgagatatttgaaaaagtagtcaggaaaattcctgattatttttttacaaaaaacgcttga 56486
57 F E I F E K V V R K I P D Y F F Y K N A *

Table 31

Query= sid|114822|lan|dp1ORF001 Phage dp1 ORF|36698-40390|2
(1230 letters)

>gi|928828 (L44593) ORF1904; putative [Lactococcus lactis phage BK5-T]
Length = 1904

Score = 427 bits (1086), Expect = e-118
Identities = 226/475 (47%), Positives = 281/475 (58%), Gaps = 45/475 (9%)

Query: 395 AESGKYIGVLNTNKKPSRLVPPDFTWIRLEGPKGDAGLPGAPGRDGVDPGKSGVGIAD 454
A+ YIG + P D+TW + +G+ G GA G+DGV GK GVGI
Sbjct: 820 ADYPSYIGQYTDYFIQYDSAKPSDYTWLSI---RGNDGKDGTGKDG---AGKDGVGIKT 873

Query: 455 TAITYAVSVSGTQEPENGWSEQVPELIKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGN 514
T ITYA+S SGT +P GW+ QVP L+KG++LWTKT W YTD S ETGYSV YI +DGN+
Sbjct: 874 TVITYALSSSGTDKPNWTGWSQVPTLVKGQYLWTKTVWITYDSSSETGYSVTYIAKDGN 933

Query: 515 GKDGIAKDGVGIAATEVMYASSPSATEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTD 574
G DGIAGKDGVGI T + YA S T APA GW++QVP VP GQ+LWT+T W YTD T
Sbjct: 934 GNDGIAGKDGVGIKTTITYAVGTSGTTAPASGWNSQVFNVPAGQFLWTKTVWITYDNTS 993

Query: 575 EIGYSVSRMGEQGPKGDAAGR---DGIAGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVP 630
E GYSV+ MG +G KGD G +GIAGK+G G+K+T+++Y SP + P G W++ VP
Sbjct: 994 ETGYSVAMMGVKGDKGDPGNGTNGIAGKDGKIKATAITYQASPNGTAPTGTWSASVP 1053

Query: 631 SLIKGQYLWTRTIWITYDSTTETGYQKTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGST 690
+ KG +LWTRTIWITYD+TTETGY Y+ +GN+G +G GKDG GIK+TTITYAGST
Sbjct: 1054 PVAKGSFLWTRTIWITYDNTTETGYAVAYMGNGNNGHDGFPKGDTGIKTTITYAGST 1113

Query: 691 SGTVAPTSNTWSAIPNVQPGPFLWTKTVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXX 750
SGT P + WTS +P V G +LWTKTVW YTD+TSETGYSV+ +G
Sbjct: 1114 SGTTTPNNGWTSTVPTVAEGNYLWTKTVWITYDNTSETGYSVAMMG-----VKGDKGDP 1167

Query: 751 XXXXXXXXXXXXADGRS-QYTHLAFSNSPNGEFSGHTDSGRAYVVGQYQDFNVHSDPAAYT 809
DG+ + T + + SPNG A G + P +K +T
Sbjct: 1168 GNGTNGIAGKDGKIKATAITYQASPNGT-----TAPTGTWSASVPPVAKGSFLWT 1219

Query: 810 WTKN-----KGNDGAQGIPGKPGADGKTNYFHIAAYASSADGS 846
T W GN+G G PGK G KT I YA S G+
Sbjct: 1220 RTIWTYDNTTETGYAVAYMGNGNNGHDGFPKGDTGIKTT--TITYAGSTSGT 1272

Score = 396 bits (1007), Expect = e-109
Identities = 208/449 (46%), Positives = 260/449 (57%), Gaps = 42/449 (9%)

Query: 421 IRLEGPKGDAGLPGAPGRDGVDPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
+ + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
Sbjct: 1155 VAMMGVKGDKG---DPGNGTNGIAGKDGKIKATAITYQASPNGTAPTGTWSASVPPV 1211

Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGN SGKDGIAKDGVGIAATEVMYASSPSA 540
KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
Sbjct: 1212 AKGSFLWTRTIWITYDNTTETGYAVAYMGNGNNGHDGFPKGDTGIKTTITYAGSTSG 1271

Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQDEIGYSVSRMGEQGPKGDAAGR---DGI 597
T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G +GI
Sbjct: 1272 TTPPNNGWTSTVPTVAEGNYLWTKTVWITYDNTSETGYSVAMMGVKGDKGDPGNGTNGI 1331

Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKQYLWTRTIWITYDSTTETGYQ 656
AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWITYD+TTETGY
Sbjct: 1332 AGKDGKIKATAITYQASPNGTAPTGTWSASVPPVAKGSFLWTRTIWITYDNTTETGYA 1391

Query: 657 KTYIPKDGNDGKNGIAGKDGVGIKSTTITYAGSTSGTVAPTSNTWSAIPNVQPGFFLWTK 716
Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK
Sbjct: 1392 VAYMGNGNNGHDGFPKGDTGIKTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1451

Query: 717 TVWNYTDDTSETGYSVSKIGETXXXXXXXXXXXXXXXXXXXXXADGRS-QYTHLAFSNS 775
TVW YTD+TSETGYSV+ +G DG+ + T + + S
Sbjct: 1452 TVWITYDNTSETGYSVAMMG-----VKGDKGDPGNGTNGIAGKDGKIKATAITYQAS 1505

399

Query: 776 PNGEGFSHTDSGRAYVQYQDFNPVHSDPAAYTWTW-----KGND 817
 PNG A G + P +K +T T W GN+
 Sbjct: 1506 PNGT-----TAPTGTWSASVPPVARGSLWTRTIWYTDNTTETGYAVAYMGNTGNN 1557

Query: 818 GAQGIPGKPGADGKTNYPHIAVASSADGS 846
 G G PGK G KT I YA S G+
 Sbjct: 1558 GHDGFPKDGDTGIKTT--TITYAGSTSGT 1584

Score = 384 bits (977), Expect = e-105
 Identities = 179/322 (55%), Positives = 222/322 (68%), Gaps = 7/322 (2%)

Query: 421 IRLEGPKGDAGLPGAPGRDGVDPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
 + + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
 Sbjct: 1311 VAMMGVKGDKG---DPGNGTNGIAGKDGKGIKATAITYQASPNGTAPTGTWSASVPPV 1367

Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGN SGKDGKGIAGKDGVGIAATEVMYASSPSA 540
 KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
 Sbjct: 1368 AKGSFLWTRTIWYTDNTTETGYAVAYMGNTGNNNGHDGFPKDGDTGIKTTTITYAGSTSG 1427

Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDGIGYSVSRMGEGQPKGDAGR---DGI 597
 T P GW++ VPTV G YLWT+T W YTD T E GYSV+ MG +G KGD G +GI
 Sbjct: 1428 TTPPNNGWTSTVPTVAEGNYLWTKTVWYTDNTSETGYSVAMMGVKGDKGDPGNGTNGI 1487

Query: 598 AGKNGIGLKSTSVSYGISPTDSAIP-GVWASQVPSLIKGQYLWTRTIWYTDSTTETGYQ 656
 AGK+G G+K+T+++Y SP + P G W++ VP + KG +LWTRTIWYTD+TTETGY
 Sbjct: 1488 AGKDGKGIKATAITYQASPNGTAPTGTWSASVPPVAKGSFLWTRTIWYTDNTTETGYA 1547

Query: 657 KTYIPKDGNDGKNGIAGKDGVGKISTTITYAGSTSGTVAPTSNWTSAIPNVQPGFFLWTK 716
 Y+ +GN+G +G GKDG GIK+TTITYAGSTSGT P + WTS +P V G +LWTK
 Sbjct: 1548 VAYMGNTGNNNGHDGFPKDGDTGIKTTTITYAGSTSGTTPPNNGWTSTVPTVAEGNYLWTK 1607

Query: 717 TVWNYTDDTSETGYSVSKIGET 738
 TVW YTD++ ETGYSV K+G T
 Sbjct: 1608 TVWAYTDNSFETGYSVKGNGNT 1629

Score = 201 bits (507), Expect = 2e-50
 Identities = 121/297 (40%), Positives = 156/297 (51%), Gaps = 19/297 (6%)

Query: 421 IRLEGPKGDAGLPGAPGRDGVDPGKSGVGIADTAITYAVSVSGTQEPENGWSEQVPEL 480
 + + G KGD G PG +G +G+ GK G GI TAITY S +GT P WS VP +
 Sbjct: 1467 VAMMGVKGDKG---DPGNGTNGIAGKDGKGIKATAITYQASPNGTAPTGTWSASVPPV 1523

Query: 481 IKGRFLWTKTFWRYTDGSHETGYSVAYIGQDGN SGKDGKGIAGKDGVGIAATEVMYASSPSA 540
 KG FLWT+T W YTD + ETGY+VAY+G +GN+G DG GKDG GI T + YA S S
 Sbjct: 1524 AKGSFLWTRTIWYTDNTTETGYAVAYMGNTGNNNGHDGFPKDGDTGIKTTTITYAGSTSG 1583

Query: 541 TEAPAGGWSTQVPTVPGGQYLWTRTRWRYTDQTDGIGYSVSRMGEGQPKGDAGRDIAGK 600
 T P GW++ VPTV G YLWT+T W YTD + E GYSV +MG GP AG +G GK
 Sbjct: 1584 TTPPNNGWTSTVPTVAEGNYLWTKTVWAYTDNSFETGYSVKGNGTGP---AGSNGNPGK 1640

Query: 601 NGIGLKSTSVSYGISPTDSAIPGVWASQVPSLIKG-QYLWTRTIWYTDSTTE--TGYQK 657
 + T+ G++ S + + ++ G +Y W W + G
 Sbjct: 1641 VVSDTEPTTKFKGLTWKYSVGVDMPLGNGTKILAGTEYWNNGNWLALYEINAHNINGDNL 1700

Query: 658 TYIPKDGNDGK-NGIAGKDGVGKISTTITYAGS-----TSGTVAPTSNWTSAIPNVQ 708
 + DGK I G +GV + T T GS +S + T N T AI N Q
 Sbjct: 1701 SVTNGTFFKDGKIESINGVNGV---NGTTIEGSHLQIHSSDSTTNTEN-TLAIDNRQ 1753

Query= sid|114823|lan|dp1ORF002 Phage dp1 ORF|32386-35835|1
 (1149 letters)

>dbj|BAA31888| (AB009866) orf 15 (bacteriophage phi PVL)
 Length = 694

Score = 280 bits (709), Expect = 3e-74
 Identities = 157/465 (33%), Positives = 257/465 (54%), Gaps = 28/465 (6%)

Query: 40 QIGSALTGLGKGLTTAVTLPLMGFAAASIKVNEFQAQMSRVQAIAGATAEELGRMKTQA 99
 +IG+++ +G+ +T VT P++ A + K G EF M +V+A +GAT EE +K +A
 Sbjct: 151 EIGNSMKNVGRNMTMYVTAPVVAGFAVAAKGIIEFDDSMRKVKATSGATGEEFEALKKKA 210

400

Query: 100 IDLGAKTAFSAKEAAQGMENLASAGFQVNEIMDAMPGVLDLXXXXXXXXXXXXXXXXXMASSL 159
++GA T FSA ++A+ + +A AG+ ++M+ + GV+DL + L
Sbjct: 211 REMGATTKFSASDSAEALNYMALAGWDSKQMMELSGVMDLAAASGEELGAVSDIVTDGL 270

Query: 160 RAFGLEANQAGHVADVFAAAADTNAETSDMAEAMKYVAPVAHSMGLSLEETAASIGIMA 219
AFGL+A +GH+ADV A+ ++ N + + EA KYVAPVA ++G ++E+T+ +IG+M+
Sbjct: 271 TAFGLKAKDSGHLADVLAQTSSKANTDVRGLGEAFKYVAPVAGALGYTTIEDTSIAIGLMS 330

Query: 220 DAGIKGSQAGTTLRGALSRIAKPTKAMVKSQMQLGVSFYDANGNMIPLEQIAQLKTATA 279
+AGIKG +AGT LR + ++ PT+AM M+ LG+S D+NG MIP+R+ + QL+
Sbjct: 331 NAGIKGEKAGTALRTMFTNLSSPTRAMGNEMERLGISITDSNGKMPMRKLLDQLREKFK 390

Query: 280 GLTQEERNRHLVTLYGQNSLSGMLALLDAGPEKLDKMTNALVNSDGAAKEMAETMQDNLA 339
L+++++ T++G+ ++SG LA+++A E K+T ++ +S GA+K MA+TM+ L
Sbjct: 391 HLSKDDQQAASATIFGKEAMSGALAIINASDEDYQKLTKSIDSSTGASKRMADTMESGLG 450

Query: 340 SKIEQMGGAFESVAIIVQILEPALAKIVGAITKVLEAFVNMSPIGQKMWVIFAGMVAAL 399
K+ + E +A+ + +EPAL IV A +KV+ + Q VV F VA L
Sbjct: 451 GKLRTRLRSQLEBALTIYDRIEPALKIIVSAPSKVVTWVTKLPTSQQLAVVGFGLFVAVL 510

Query: 400 GPLLLIAGM-----VMTTIVKLRIAIQFLGPAFMGTMGTIAGVIAIF----- 441
GPL+ + G+ MT + L I + F IA ++ +F
Sbjct: 511 GPLVFMFLFISVMGNAMTVLGPLLINVNKASGLFAFLRTKIASLVKLPFILGVSISSLT 570

Query: 442 -----YALVAV---FMIAYTKSERFRNFINS LAPAIKAGFGGA 476
ALV + F AY +SE FRN +N + F A
Sbjct: 571 LPITLIVGALVGIGIAFYQAYKRSETFRNIVNQAISGVANAFKAA 615

Query= sid|114824|lan|dp1ORF003 Phage dp1 ORF|53538-55877|3
(779 letters)

>sp|P43741|DPO1_HABIN DNA POLYMERASE I (POL I) >gi|1074025|pir|E64098 DNA polymerase I
(polA) homolog - Haemophilus influenzae (strain Rd KW20)
>gi|1573871 (U32767) DNA polymerase I (polA)
[Haemophilus influenzae Rd]
Length = 930

Score = 191 bits (481), Expect = 1e-47
Identities = 148/553 (26%), Positives = 262/553 (46%), Gaps = 60/553 (10%)

Query: 63 RLELITEEAKLEQYVDKMIEDGIGSIDVETDGLDTHDELAGVCLYSPSQKGIYAPVNVH 122
+ E + +A L ++++K+ + ++D ETD LD + L G+ + + Y P+
Sbjct: 333 KYETILLTQADLTRWIEKLNAAKLIADVOTETDSDLYMSANLVGISFALENCEAAYLPLQLD 392

Query: 123 SNMTKMRIKNQISPEFMKKMLQRIVDSGIPVIYHNSKFDMSIYWRVGVMNEPAWDTYL 182
++ + +K +L+ + I I N KFD +SI+ R G+++ +DT L
Sbjct: 393 YLDAPKTLEKSTALAAIKPILE---NPNHKGIGQNIKFD-ESIFARHGIELQGVEFDTML 448

Query: 183 AAMLNENESHSLKSLHXYVRNEENAFAKFNDFKGIFFSLIPPDVAYMYAAYDPLQT 242
+ LN H++ L +Y+ +E A + + F+ IP + A YAA D T
Sbjct: 449 LSYTLNSTGRHNMDLAKRYLGHETIAPESLAGKGSQTLTFNQIPLEQATEYAAEDADVT 508

Query: 243 FELYEFQEYQLTPGTEQCEEYNLEKVSFVNLHNIEMPLIKVLFDMVEYGVLDLQDKLAEIR 302
+L + L E Y +E+PL+ VL ME GV +D D L
Sbjct: 509 MKLQALWLKLEQEPTLVELYK-----TMELPLLHVLNRMERTGVLDSDALFMQS 559

Query: 303 EQPTANMNEAEQEQQLVSEWQPEIEELRQTNFQSYQKLEMDARGRVTVSISSTQLAIL 362
+ + + E++ L + +++S QL +
Sbjct: 560 NEIASRLTALEKQAYALAGQ-----PFNLASTKQLQEI 592

Query: 363 FYDIMGKSPERDKPRG---TGESIVEH--FDNDISXXXXXXXXXXXXVSTYTT-LDQHL 416
+D + L ++ P+G T E ++E + +++ STYT L Q +
Sbjct: 593 LFDKLELPVLQKT- PKGAPSTNEEVLELSYSHLKPILVKHRLSKLSTYTDKLPQMV 651

Query: 417 AKPDNRHITTFKQYGAKTGRMSENPNLQNIQPSRGE-GAVVRQIFAASEGHYIIGSDYSQ 475
R+HT++ Q TGR+SS +PNLQNIQ R E G +RQ F A EG+ I+ +DYSQ
Sbjct: 652 NSQTGRVHTSYHQAVTATGRSSDPNLQNIQIRNEEGHRIHQAFIAREGYSIVAADYSQ 711

Query: 476 QEPRSLAELSGDESMRHAYEQNLDLYSVIGSKLYGVFYEBCLEFPYDGTNKEGKLRRNS 535
E R +A LSGD+ + +A+ Q D++ ++++GV +E T+++ R +
Sbjct: 712 IELRLMAHLSGDQGLINAFSQGKDIHRSTAEIFGVSLDE-----VTSEQ-----RRN 759

Query: 536 VKSVLLGLMYGRGANSIAEQMNVSVKEANKVIEDFFTEFPKVDYIIFVQQQAQDLGYVQ 595
K++ GL+YG A ++ Q+ +S +A K ++ +F +P V ++ +++++A+ GYV+

401

Sbjct: 760 AKAINFGLIYGMSAFGLSRQLGISRADAQKYMPLYFQRYPSVQQFMTDIREKAKAQGYVE 819

Query: 596 TATGRRRLPDMS 608

T GRR LPD++

Sbjct: 820 TLFGRRLLYLPDIN 832

Score = 46.9 bits (109), Expect = 5e-04

Identities = 34/123 (27%), Positives = 66/123 (53%), Gaps = 16/123 (13%)

Query: 663 EIKDQAKAEGI-----LIKDNGGKIADAQRQCLNSVIQGTAAADMTKYAMIKV 709
+I+++AKA+G + N + A+R +N+ +QGTAA+ K AMIK+

Sbjct: 807 DIREKAKAQGYVETLFGRRLLYLPDINSSNAMRRKGAERVAINAPMQGTAAIIKRAMIKL 866

Query: 710 HNDDELKELGFLMIPVHDELLGEVPIKNAKGAERLTEVMIEAAKDIISLPMKCDPSIV 769

++ + +++ VHDEL+ EV + E++ + M EAA +++ +P+ + +

Sbjct: 867 -DEVIRHDPDIEMIMQVHDELVEVRSEKVAFFREQIKQHM-EAAAEV-VPLIVEVGVG 923

Query: 770 ERW 772

+ W

Sbjct: 924 QNW 926

Query= sid|114825|lan|dp1ORF004 Phage dp1 ORF|40401-42440|3
(679 letters)

>emb|CAB07981| (Z93946) hypothetical protein [bacteriophage Dp-1]
Length = 532

Score = 1011 bits (2585), Expect = 0.0

Identities = 497/499 (99%), Positives = 498/499 (99%)

Query: 1 MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLN 60
MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLN

Sbjct: 1 MTKFINSYGPLHLNLYVEQVSQDVTNNSSRVSWRATVDRDGAYRTWTYGNISNLSVWLN 60

Query: 61 SSVHSSHPDYDTSGEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL 120

SSVHSSHPDYDTSGEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL

Sbjct: 61 SSVHSSHPDYDTSGEVTLASGEVTVPHNSDGTKTMSVWASFDPNNGVHGNITISTNYTL 120

Query: 121 DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGNHTTSVSFT 180

DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGNHTTSVSFT

Sbjct: 121 DSIPRSTQISSFEGNRNLGSLHTVIFNRKVNSFTHQVWYRVFGSDWIDLGNHTTSVSFT 180

Query: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT 240

PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT

Sbjct: 181 PSLDLARYLPKSSSGTMDICIRTYNGTTQIGSDVYSNGWRFNIPDSVRPTFSGISLVDTT 240

Query: 241 SAVRQILTGNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF 300

SAVRQILTGNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF

Sbjct: 241 SAVRQILTGNFLQIMSNIQVNFNNASGAYGSTIQAFHAELVGKNQAINENGGKLGMMNF 300

Query: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTQRNPAAIQALRNAKVAPI 360

NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTQRNPAAIQALRNAKVAPI

Sbjct: 301 NGSATVRAWVTDTRGKQSNVQDVSINVIEYYGPSINFSVQRTQRNPAAIQALRNAKVAPI 360

Query: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISLMTNSSANLAGNYGPKSYIV 420

TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISL+TNSSANLAGNYGPKSYIV

Sbjct: 361 TVGGQQKNIMQITFSVAPLNTTNFTEDRGSASGTFTTISLMTNSSANLAGNYGPKSYIV 420

Query: 421 KAKIQDRFTSTEFSAATVATESVVLNYDKDGRGLGVGKVVEQKAGSIDAAGDIYAGGRQVQ 480

KAKIQDRFTSTEFSAATV TESVVLNYDKDGRGLGVGKVVEQKAGSIDAAGDIYAGGRQVQ

Sbjct: 421 KAKIQDRFTSTEFSAATVPTESVVLNYDKDGRGLGVGKVVEQKAGSIDAAGDIYAGGRQVQ 480

Query: 481 QFQLTDNNGALNRGQYNDV 499

QFQLTDNNGALNRGQYNDV

Sbjct: 481 QFQLTDNNGALNRGQYNDV 499

Query= sid|114827|lan|dp1ORF006 Phage dp1 ORF|45296-46987|2
(563 letters)

>gb|AAD18987| (AE001666) SWI/SNF family helicase_2 [Chlamydia pneumoniae]
Length = 1166

Score = 171 bits (429), Expect = 1e-41

Identities = 150/522 (28%), Positives = 254/522 (47%), Gaps = 55/522 (10%)

402

Query: 46 SSNNFE-LPYKYFNNVIDALDEWELHIFGELDKVDQYIDSRNRIASSSNEQFSFKTTPF 104
S + FE LP + ++ + L E + I GE++ D QD + T

Sbjct: 659 SLDQFEALPVNF--SMSERLIEIQKQIRGEIEFDFQD-----VPQQIQATLRSYQTEG 709

Query: 105 AHQVECFEYAEHPCFLLGDEQGLGKTKQAIDIAVSRKASFH--CLIVCCISGLKWNWA 162
H +E + H +L D+ GLGKT QAI IAV++ K C ++ C + L +NW

Sbjct: 710 VHWLE--RLRKMHLNGILADDMGLGKTLQAI-IAVTQSKLEKSGCSLIVCPTSLVYNWK 766

Query: 163 KEVGIHSNESAHILGSRVTGDKGLVIDGV-SKRAEDLLGGHDEFFLITNIETLRDAVFIK 221
+E + E LVIDGV S+R + L D IT+ L+ V

Sbjct: 767 EEFRKFNPEFR-----TLVIDGVPSQRRKQLTALADRDVAITSYNLLQKDV--- 812

Query: 222 YLNELTKSGEIGMVIIDEIHKCKNPSSKQASIQKLQSYKMGTLGTPLMNPIDVFNVM 281
EL KS V++DE H KN +++ S++ +QS +++ LTGTP+ N+ +++++

Sbjct: 813 ---ELYKSFRFDYVVLDEAHHIKNRTTRNAKSVKMIQSDHRLILTGTPIENSLEELWSLF 869

Query: 282 KNLGAEHHTLTQFKERYCIVDQFNQITGYR-----NLAEELRELVDYMLRRTKEEVL-DL 335
+L L +R+ V ++ + Y N+ L++ V+ ++LRR KE+VL DL

Sbjct: 870 DFLMPG---LLSSYDRF--VGKYIRTGNMGNKADNMVALKKVSPFILRMKEDVLKOL 924

Query: 336 PEKIRVTEYVDMNSKQSKIY-----KEVLTCLVQEIDKVKLMPNPLAETIRLRQATGN 388
P + + + Q ++Y K+ L++LV++ ++ + LA RL+Q +

Sbjct: 925 PPVSEILYHCHLTESQKELYQSYAASAKQELSRVLKQEGFERIHIHVLATLTLKQICCH 984

Query: 389 PSILTTQDVK---SCKFERCIEIVEECIQQKSCVIFSNEKVEIPLAKIL-SKTVKCNL 444
P+I + S K++ +++++ + G V+FS + K++ + K L S+ +

Sbjct: 985 PAIFAKDAPEPGDSAKYDMLMDLLSSLVDSGHKTVVFSQYTKMLGIKKDLESRGIPFVY 1044

Query: 445 VTGETADKFNEIEEPMNHRKASVILGTIGALGTGFTLTAKDTVIFLDSPTRAEKDQAE 504
+ G T ++ + + +F V L ++ A GTG L ADTVI D W A ++QA D

Sbjct: 1045 LDGSTKNRLDLVNQFNEDPSLLVFLISLKAGTGLNLVGADTVIHYDMWNPAPVENQATD 1104

Query: 505 RCHRIGAKSSVTIYTLVAKGTVDERIEDLIERKGELADYIVD 546
R HRIG SV+ Y LV T++E+I L RK L +++

Sbjct: 1105 RVHRIGQSRVSYSYKLVTLNTEIEKILTQNRKKSLLVKKVIN 1146

Query= sid|114828|lan|dp1ORF007 Phage dp1 ORF|22230-23621|3
(463 letters)

>gi|2444105 (U88974) ORF26 [Streptococcus thermophilus temperate bacteriophage
O1205]
Length = 411

Score = 88.9 bits (217), Expect = 7e-17
Identities = 80/315 (25%), Positives = 133/315 (41%), Gaps = 48/315 (15%)

Query: 139 QGVTLAGIFCDEVALMPESFVNQATGRCSVTGSKMWFSCNPANPNHYFKKNWIDKQVEKR 198
+G T G + +E +L E + RCS G+++ + NP NPNH+ +++I K + +

Sbjct: 121 RGFTAFGAYVNEASLANELVFKEIISRCSGDGRVVDNSNPDNPNHNLNRDYIGKN-DGK 179

Query: 199 ILYLHFTMDNPSLT---DSIKRREYKMYAGVFRKRFILGLWVTADGLVYSMFNEEQHV 254
I+ F +DDN L+ DSIK K G F R ILGLW A+G +Y+ ++ + HV

Sbjct: 180 IIDFSFKLDNTPLSKRYIDSIAATPK---GKPYDRDILGLWTVAGAIYADYDSKIHV 236

Query: 255 KKLNIIEFDRLFVAGDFGIYNATTFGLYGFSKRHKRYHLIESYHSGREAEQLTEADVNS 314
E R F D+G + + + G ++L++ +E + + +A

Sbjct: 237 VDLEPEMKRYFGGIDWGYTHYGSIVIVG-EGVDNNFYLVDGVAAQFKEIDWVVEQA---- 291

Query: 315 NIQFSSVLQKTTKEYANDLVDMIRGKQIEYIILDPSASAMIVELQKHPYIAR---KNIP 371
+K T Y N + + ++AR + I

Sbjct: 292 -----RKLGTGIYGN-----IPFYADSARPEHVARFENEGFDI 323

Query: 372 IPARNDVTLGISFHAELLAENRFTLDPSNT-HDIDEYYAYSWDSKASQTGEDRVIKEDH 430
+ A V GI A+L E + + DE Y Y W ++ +D +KE D

Sbjct: 324 MNANKSVIAGIELIAKLFKEKLYVKRGFVPRFFDEIYQYRWKENST---KDEPLKEFDD 380

Query: 431 CMDRNRYACLTDALI 445
+D RYA +D +I

Sbjct: 381 VLDSVRYAIYSDYVI 395

Query= sid|114829|lan|dp1ORF008 Phage dp1 ORF|49624-50961|1
(445 letters)

>gb|AAD19901| (AF100420) DnaB replication fork helicase [Thermus aquaticus]

403

Length = 444

Score = 67.5 bits (162), Expect = 2e-10
 Identities = 69/248 (27%), Positives = 111/248 (43%), Gaps = 14/248 (5%)

Query: 147 GERLGISTGFEXXXXXXXXXXXXXXXXXXIVIMARPGQKS-WTIDKMLATAWKNHVDVLLYS 205
 GE G+ TGF+ I I ARP GK+ + + A K G V +YS
 Sbjct: 178 GEVAGVRTGFKELDQLIGTLGPGSLNI-IAARPAMGKTAFALTIAQNAALKEGVGVGIYS 236

Query: 206 GEMSEMQVGARIDTILSNVSINSITKGIWNDHQFEKYEDHIQAMTEAENSLVVVTPFMIG 265
 EM Q+ R+ + +N+ G D F + D ++EA + TP +
 Sbjct: 237 LEMPAAQLTLRMCMSEARIDMNRVRLGQLTDRDFSRLVDVASRLSEAP-IYIDTDPDLTL 295

Query: 266 GKNLTPAILDSMISKYRPSVVGIDQLSLMS--ESYPSREQKRIQYANITMDLYKISAKYG 323
 + A ++S+ + ++ ID L LMS S S E ++ + A I+ L ++ + G
 Sbjct: 296 ME--VRARARRLVSQNQVGLI IIDYLQLMSGPGSGKSGENRQOEIAAISRLKALARELG 353

Query: 324 IPIVLNVQAGRSKTEGAESMELEHIAESDGVGNASRVIAMKRD-----EKSGILEL 376
 IPI+ Q R+ + + L + ES + Q+A V+ + RD EK+GI E+
 Sbjct: 354 IPIIALSQLSRAVEARPKNRPMLSDLRESGSIEQDADLVMPFIYRDEYYNPHSEKAGIAEI 413

Query: 377 SVVKNRYG 384
 V K R G
 Sbjct: 414 IVGKQRNG 421

Query= sid|114831|lan|dp1ORF010 Phage dp1 ORF|8699-9859|2
 (386 letters)

>gi|2760912 (AF037258) RecA protein [Chlorobium tepidum]
 Length = 346

Score = 133 bits (331), Expect = 2e-30
 Identities = 99/340 (29%), Positives = 164/340 (48%), Gaps = 66/340 (19%)

Query: 44 GGLPRKRVVEFFGPESGKTTSDIVKNAQMVFXXXXXXXXXXXXXXXXXXNARASKASKT 103
 GGLPR RV E +GPESGKTT AL + AQ
 Sbjct: 67 GGLPRGRVTEIYGPESGKTTLALHAIIEAQ-----KNG 100

Query: 104 AVKELEMQLDSLQEPKIVYLDLENTLDTWAKKIGVDVDNIWIVRPEMNSAEELQYVL 163
 + L +D E+ D +A+K+GVD++ + +PE S E+ L V
 Sbjct: 101 GIAAL-----VDAEHAFDPTYARKLGVDINALLVSQPE--SGEQALSIVE 143

Query: 164 DIFETGEVGLVVLDSLPMVSNLIDEELTKKAYAGISAPLTFESRKVTPLLTRYNAIFL 223
 + +G V ++V+DS+ +V Q ++ E+ + +++ RK+T +++ ++ L
 Sbjct: 144 TLVRSGAVDIIVDSVAALVPQAELEGEMGDSVVGQLARLMSQALRLTGAISKSSSVCL 203

Query: 224 GINQIREDMNSQYNA-YSTPGGKMKHACAVRLKFRKGDYLDENGASLTRTARNPAGNVV 282
 INQ+R+ + Y + +T GGK K +VRL RK + ++G L GN
 Sbjct: 204 FINQLRDKIGVMYGPSTTTGGKALKFYSSVRLDIRKIAQI-KDGEELV-----GNRT 255

Query: 283 ESFVEKTKAFKPRKLVSYTSLYHGIQIENDLVDVAVEFGVIQKAGAWFSIVDLETGEI 342
 + V K K P K + + Y +GI + +L+D+AVEFG+I+K+GAWFS + G
 Sbjct: 256 KVKVVKVKV-APPFKTAEFDILYEGISVLGELIDLAVEFGIHKSGAWFSYGTEKLG-- 312

Query: 343 MTDEDEEPLKFQGKANLVRRFKEDDYLFDMVMTAVHEIIT 382
 QG+ N+ + KED+ L + + V +++T
 Sbjct: 313 -----QGRENVKLLKEDETLENTIRQQVRDMLT 341

Query= sid|114832|lan|dp1ORF011 Phage dp1 ORF|28017-29096|3
 (359 letters)

>gi|2444110 (U88974) ORF31 [Streptococcus thermophilus temperate bacteriophage
 01205]
 Length = 348

Score = 187 bits (469), Expect = 1e-46
 Identities = 118/358 (32%), Positives = 187/358 (51%), Gaps = 21/358 (5%)

Query: 3 IYDYINAGEIASYIQALPSNALQYLGPFLFNAQQTGTDISWLKGANNLPVTIQPSNYDA 62
 IYD + A IA Y AL N LG ++FP +Q GT +S++KGA+ V ++ + +D
 Sbjct: 4 IYDKVTASNIAGYFNALQENVSSTLGESIFPARKQLGTLKLSYIKGASGQSVALKAAAFDT 63

Query: 63 KASLRERAGFSKQATEMAFFRESMRLGEKDRQNLQMLLNQSSA-LAQPLITQLYNDTKNL 121

404

+R+R +M FF+E+M + E DRQ L ++ + +A L ++ ++ND L
 Sbjct: 64 NVTIRDRVSAEMHDEQMPFFKEAMLVKENDRQQLNLVKDSGNAVLVNTIVAGIFNDNLTL 123
 Query: 122 VDGVEAQAEYMRMQLLQYQKFTVKSTNSEAQYTYDYNMDAQYAVTKKWTNPAESDPIA 181
 V+G A+ E MRMQ+L GK S Y D K+Q V+K W P + P+A
 Sbjct: 124 VNGARARLEAMRMQVLATGKIAFTSDGVNKKIDYGVKPDHKKQ--VSKSWAEPG-ATPLA 180
 Query: 182 DILAAMDDIENRTGVRPTRMVLNRNTYNQMTKSDSIKKAL-AIGVQGSWENFLLLASDAE 240
 D+ A+ + G+ P R V+N T+ + K+ S K + + GS + ++ E
 Sbjct: 181 DLEDAI-ETARELGSLNPERAVMNAKTFGLIRKAASVTKVIKPLAGDGS----AVTKAELE 235
 Query: 241 KFIAEKTGLQIAVYSKKIAQFADADKLPDVGNIQFNLIDDGKVVLLPPDAVGHTWYGT 300
 +IA+ G+ I + + D G + +F DG + L+P +G+T +GTT
 Sbjct: 236 NYIADNFGVSVILENGTYRN-----DKGEVSKF--YPDGHLTLIPNGPLGNTVFGTT 285
 Query: 301 PEAFLASGGT-DAQVQLSGGPTVTITYLEKHPVNIATVVSAMVIPSFEGIDYVGVLT 357
 PE DL + T +A+V+++ G VTT PVN+ T VS V +PSFE +D V +LT
 Sbjct: 286 PEESDLFADNTVNAEVEIVDNGIAVTTTKTTDPVNVQTKVSMVALPSFERLDDVYMLT 343

Query= sid|114834|lan|dp1ORF013 Phage dp1 ORF|10215-11240|3
 (341 letters)

>sp|P09122|DP3X_BACSU DNA POLYMERASE III SUBUNITS GAMMA AND TAU
 Length = 563

Score = 182 bits (458), Expect = 2e-45
 Identities = 118/353 (33%), Positives = 176/353 (49%), Gaps = 31/353 (8%)

Query: 7 YRPQTFEEVVAQEYVKEILLNLQNGAIGHYLFXXXXXXXXXXXXXRIFAKDVN----- 60
 +RPQ FE+VV QE++ + L N L H YLF +IFAK VN
 Sbjct: 10 FRPQRFEDVVGQEHITKTQLNALLQKKFSHAYLFSGPRGTGKTSAAKIFAKAVNCEHAPV 69
 Query: 61 -----KGL-----GSPIEIDAASNNGVENVRNIIEDSRYKSMDSSEFKVYIIDEVH 105
 KG+ IEIDAASNNGV+ +R+I + ++ +KVYIIDEVH
 Sbjct: 70 DEPCNECAACKGITNGSISDVIEIDAASNNGVDEIRDIRDVKVFPASVATYKVYIIDEVH 129
 Query: 106 MLSTGAFNALLKLTLEEPSGTVFILCTTDPQKIPDTILSRVQRFDFTRIDNDDIVNQLQF 165
 MLS GAFNALLKLTLEEP +FIL TT+P KIP TI+SR QRDF RI + IV ++
 Sbjct: 130 MLSIGAFNALLKLTLEEPPEHCIFILATTEPHKIPLTIISRCQRFDFKRITSQAIVGRMNK 189
 Query: 166 IIESENEEGAGYSYERDALSFICKLANGGMRDSITRLEKVLDSHHVDMBAVSNAL---G 222
 I+++E E +L I A+GGMRD+++ L+++ +S D+ V +AL G
 Sbjct: 190 IVDAEQ-----LQVEEGSLEIIASAAGGMRDALSLLDQAISFSG--DILKVEDALLITG 242
 Query: 223 VPDYETFASLVEAIIANYDGSKCLEIVNDFHYSKDLKLVTNRNFTDFLLEVCKYWLVRDIS 282
 L +++ + + S LE +N+ GKD + + + ++ Y +
 Sbjct: 243 AVSQLYIGKLAKSLHDKNVSDALETNLNLLQQKDPAKLIEDMIFYFRDMLLYKTAPGLE 302
 Query: 283 ITQLPAHFESKLEQFCEAFQYPTLLWMLEEMNELAGVVKWEPAKPIIETKLL 335
 + + E L M++ +N+ +KW + + E ++
 Sbjct: 303 GVLEKVKVDETFRELSEQIPAQALYEMIDILNKSHEMKWTNHPRIFFEVAVV 355

Query= sid|114835|lan|dp1ORF014 Phage dp1 ORF|50961-51974|3
 (337 letters)

>sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64227 DNA primase (dnaE) homolog
 MG250 - Mycoplasma genitalium (SGC3) >gi|3844848
 (U39704) DNA primase (dnaE) [Mycoplasma genitalium]
 Length = 607

Score = 57.0 bits (135), Expect = 2e-07
 Identities = 53/190 (27%), Positives = 89/190 (45%), Gaps = 17/190 (8%)

Query: 146 EELDKYRFIHP-----YMYERKLDELIEFMFDVGYDK--LHDCITPPVRNLKGETVFF 196
 E +++Y FI+P Y++ K + + FD K + I P+ + G V F
 Sbjct: 170 ESMERYPFINPKIKPSELYLFS-KTNQQLGFFDFNTKKATFQNMIPIHDFNGNPVGF 228
 Query: 197 NRRSVRSKFHQYGEDDPKTEFLYGQYELVAFRDYFEKPISQVFVTESVINCLTLWSMKIP 256
 + RSV + ++ EF + + EL+ K ++Q+F+ E + TL + K
 Sbjct: 229 SARSDVNINKLKYKNSADHEF-FKKGELLFNHRLNKNLNLQFLIVEGYFDVFTLTNSKFE 287
 Query: 257 AVALMGVGGGN-QINLLKR--LPYRNIVLALDPDNAGQTAQEKLYRQLKRSK-VVRFLNY 312
 AVALMG+ + QI +K + +VLALD D +GQ A L +L + +V + +
 Sbjct: 288 AVALMGLALNDVQIKAIKAHFKELQTLVLALDNDASGQNAVFSLEIKLNNNNFIVEIVQW 347

Query: 313 PKEFYDNKWD 322
 + D WD
 Sbjct: 348 EHNKYD--WD 355

Query= sid|114837|lan|dp1ORF016 Phage dpl ORF|43413-44303|3
 (296 letters)

>emb|CAB07986| (Z93946) N-acetylmuramoyl-L-alanine amidase [bacteriophage Dp-1]
 Length = 296

Score = 661 bits (1686), Expect = 0.0
 Identities = 296/296 (100%), Positives = 296/296 (100%)

Query: 1 MGVDIEKGVAMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60
 MGVDIEKGVAMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH
 Sbjct: 1 MGVDIEKGVAMQARKGRVSYSMDFRDGPDSYDCSSSMYYALRSAGASSAGWAVNTEYMH 60

Query: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSNDIIHCNYAYDGIS 120
 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSNDIIHCNYAYDGIS
 Sbjct: 61 AWLIENGYELISENAPWDAKRGDIFIWGRKGASAGAGGHTGMFIDSNDIIHCNYAYDGIS 120

Query: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180
 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE
 Sbjct: 121 VNDHDERWYYAGQPYYYVYRLTNANAQPAEKKLGWQKDATGFWYARANGTYPKDEFEYIE 180

Query: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYFNRDGSMTGW 240
 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYFNRDGSMTGW
 Sbjct: 181 ENKSWFYFDDQGYMLAEKWLKHTDGNWYWFDRDGYMATSWKRIGESWYFNRDGSMTGW 240

Query: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296
 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV
 Sbjct: 241 IKYYDNWYYCDATNGDMKSNAFIRYNDGWYLLLPDGRLADKPQFTVEPDGLITAKV 296

Query= sid|114841|lan|dp1ORF020 Phage dpl ORF|1864-2658|1
 (264 letters)

>emb|CAB13247| (Z99111) similar to coenzyme PQQ synthesis [Bacillus subtilis]
 Length = 243

Score = 217 bits (548), Expect = 5e-56
 Identities = 117/248 (47%), Positives = 163/248 (65%), Gaps = 15/248 (6%)

Query: 23 MPIMEIFGPTTIQEGMVGQKTIFIRTGGCDYHCNWCDSAFTWNGTTEPE--YITGKEAA 80
 +P++EIFGPTTIQEGMVGQKT+F+RT GCDY C+WCDSAFTW+G+ + + ++T +E
 Sbjct: 5 IPVLEIFGPTTIQEGMVGQKTMFVRTAGCDYSCSWCDSAFTWDGSAKKDIRWMTAEEIF 64

Query: 81 SRILKLAFNDKGEQICNHVTLTGGNPALINEPMAKMISILKEHGFKFLETQGTFRFQWF 140
 + + D G +HVT++GGNPAL+ + + I +LKE+ + LETQGT +Q+WF
 Sbjct: 65 AEL-----KDIGGDAFVSHVTISGGNPALLKQ-LDAFIELLKENNIRAALETQGT VYQDWF 118

Query: 141 KEVSDITISPKPPSSGMRTNMKILEAIVDRM--NDENLDWSFKIVIPDENLAYARDMFK 198
 + D+TISPKPPSS M TN + L+ I+ + ND S K+VIF++ DL +A+ + K
 Sbjct: 119 TLIDDLTISPKPPSSKMVTNFQKLDHILTSQENDRQHAVSLKVIFNDEDELEFARTVHK 178

Query: 199 TFEGKLRPVNYSVGANANAY--EEGKISDRLLLEKLGWLWDKVYEDPAFNNVRPLPQLHTL 256
 + G YL VGN + + ++ + LL K L DKV D N VR LPQLHTL
 Sbjct: 179 RYPG---IPFYLQVGNDDVHTTDDQSLIAHLLGKYEALVDKVAVDALNLVRVLPQLHTL 235

Query: 257 VYDNKRGV 264
 ++ NKRGV
 Sbjct: 236 LWGNKRGV 243

406

Query= sid|114842|lan|dp1ORF021 Phage dp1 ORF|2504-3295|2
(263 letters)

>sp|P19465|GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >gi|98411|pir||A38256 GTP
cyclohydrolase I (EC 3.5.4.16) - Bacillus subtilis
>gi|143231 (M37320) regulatory protein [Bacillus
subtilis] >gi|143799 (M80245) MtrA [Bacillus subtilis]
>gi|2634696|emb|CAB14194| (Z99115) GTP cyclohydrolase I
[Bacillus subtilis]
Length = 190

Score = 208 bits (523), Expect = 4e-53
Identities = 103/185 (55%), Positives = 133/185 (71%), Gaps = 1/185 (0%)

Query: 80 VTLDNTEAAVQRLFGLLGEDAERDGLQDTPFRFVKALAEHTVGYREDPKLHLEKTFDVDH 139
V + E AV+++ +GED R+GL DTP R K AE G EDPK H + F +H
Sbjct: 4 VNKEQIEQAVRQILEAIGEDPNREGLLDTPKRVAKMYAEVFSGLNEDPKHFQTIFGENH 63

Query: 140 EDLVLVKDI PFNSLCCEHHLAPFVGKVHIAIYIPKD-KITGLSKFGRVVEGYAKRLQVQERL 198
E+LVLVKDI F+S+CEHHL PF GK H+AYIP+ K+TGLSK R VE AKR Q+QER+
Sbjct: 64 EELVLVKDIAFHSCEHHLVFPYGAHVAYIPRGKVTGLSKLARAVEAVAKRPQLQERI 123

Query: 199 TQIADAIQEVLNPQAVAVIVEAEHTCMSEGRGIKKHGATTVTSTMRGLFQDDASARAELL 258
T IA+I E L+P V V+VEAEH CM+ RG++K GA TVTS +RG+F+DDA+ARAE+L
Sbjct: 124 TSTIAESIVETLDPHGVMVVVEAEHMCMTMRGVRKPGAKTTSVAVRGVFKDDAAARAELV 183

Query: 259 QLIKK 263
+ IK+
Sbjct: 184 EHIKR 188

Query= sid|114843|lan|dp1ORF022 Phage dp1 ORF|30896-31675|2
(259 letters)

>gi|2347102 (U77367) internalin [Listeria monocytogenes]
Length = 821

Score = 55.0 bits (130), Expect = 5e-07
Identities = 44/149 (29%), Positives = 63/149 (41%), Gaps = 13/149 (8%)

Query: 119 FRMNIYVPNYVG--DSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPV 176
F + VFN + D + + NN T AP L Y PE +K + K +
Sbjct: 383 FSKTSLVPNNITSIDGTLIAPETISNNGTYDAPNLKWSLPNYLPE--VKYTFQSQKIPIGT 440

Query: 177 KSM DYVAQLPAVLR-----RVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW 231
+ +Y + L+ +VTF++ G T + V E + P+P PT G F GW
Sbjct: 441 GTSNYSFGFITQPLKELLDYKVTFNVEGNTSEVETVTEE---NLIPEPTSPKQGYTFD GW 497

Query: 232 -KVEGESTIWDFFDNHMPDRDVKLVAQFA 259
E T WDF MP D+ L A F+
Sbjct: 498 YDAETGGTKWDFTTGQMPANDLTLYAHFS 526

Score = 43.4 bits (100), Expect = 0.002
Identities = 47/195 (24%), Positives = 73/195 (37%), Gaps = 12/195 (6%)

Query: 72 YDLTFKDNFTDPEIMALIEGGTVRQGGTIAGYDT-PMLAQGASNMKPFMRNIYVPNY-- 128
YD + T + +G + GG + T M A + F +N Y N+
Sbjct: 547 YDALLNEPTTPTKQGYTFDGYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDI 606

Query: 129 ---VGDSIVNYVKITLNNCTGKAPGLSIGKEFYAPEFNIKAREATKAGLPVKSM DYVAQL 185
V + +Y + T G + + A K TK +P + A
Sbjct: 607 DGEVKNETIAYDTLLNEPTTPTKQGYTFDGYDAETGGTKWDFKTKKE-MPANDVTLYAHF 665

Query: 186 PAVLRRVTFDLNGGTGTADAVRVEAGKKISPKPVDPTLTGKAFKGW-KVEGESTIWDFFDN 244
+ FD++G T + V +A + P+P P+ TG +GW E T WDF
Sbjct: 666 TINNYQANFDIDGAV-TEEVVNYDA---LIPEPTSPSKTGFTLEGWYDAEVGGTKWDFKT 721

Query: 245 HMPDRDVKLVAQFA 259
MP D+ L A F+
Sbjct: 722 MKMPANDITLYAHFS 736

Score = 38.3 bits (87), Expect = 0.057
Identities = 42/169 (24%), Positives = 59/169 (34%), Gaps = 10/169 (5%)

407

Query: 96 QQGTTIAGYDT-PMLAQGASNMKPFMRNIYVPNYVGDSIVNYVKIT----LNNCTGKAPG 150
 + GGT + T M A + F +N Y N+ D +V + LN T
 Sbjct: 501 ETGGTKWDFTTGQMPANDLTLYAHFSVNSYQANFDIDGVVTNEAVVYDALLNEPTTPTKQ 560

Query: 151 LSIGKEFYAPEFNIKAREATKAGLPVKSM DYVAQLPAVLRRVTFDLNGGTGTADAVRVEA 210
 +Y E + +P + + A + FD++G A
 Sbjct: 561 GYTFDGWYDAETGGNKWDFKTMKMPANDVAFYAHFTINNYQANFDIDGEVKNETI----A 616

Query: 211 GKKISPKPVDPTLTGKAFKGW-KVEGESTIWD FDNHMPDRDVKLVAQF 258
 + +P PT G F GW E T WDF MP DV L A F
 Sbjct: 617 YDTLLNEPTTPTKQGYTFDGWYDAETGGTKWDFKTKEMPANDVTLYAHF 665

Query= sid|114850|lan|dp1ORF029 Phage dp1 ORF|662-1348|2
 (228 letters)

>gi|2650185 (AE001074) succinoglycan biosynthesis regulator (exsB)
 [Archaeoglobus fulgidus]
 Length = 239

Score = 119 bits (295), Expect = 2e-26
 Identities = 79/224 (35%), Positives = 113/224 (50%), Gaps = 11/224 (4%)

Query: 1 MKSVLLSGGVSATCLAEVDKMGSKNVHAIAPNYGQKHEAELENAANVAMFYGVKPTI 60
 MK+V+LLSGG+DS+T L +D G VHA+ F YGQKH E+E+A VA V+
 Sbjct: 1 MKAVMLLSGGIDSSTLLYYLLD--GGYEVHALTFYGGQKHSKEIESAEKVAKAAKVRHLK 58

Query: 61 LEIDSKIYXXXXXXLLQKGGEISHGKSYAEILAEKEVVDTVYVFRNGLMLSQXXXXXXX 120
 ++I S I+ L G+ E+ Y+E + + T VP RN ++LS
 Sbjct: 59 VDI-STIHDLISYGALTGEEVPKA-FYSEEVQRR----TIVPNRNMILLS--IAAGYAV 110

Query: 121 XXXXXXXXXXXXXXXXXXXXPDCTPEFYNSMSNAMEYGT-GGKVTLVAPLLTLTKAQVVKW 179
 PDC EF ++ A+ V + AP + +TKA +V+
 Sbjct: 111 KIGAKEVHYAAHLSDYSIYPDCRKEFVKALDTAVYLANIWTPEVVRAPFVDMTKADIVRL 170

Query: 180 GIDLDPVYPFLTRSCYESDAESCGTCATCIDRKKAFEENGMTDPI 223
 G+ L VPY LT SCYE C +C TC++R +AF NG+ DP+
 Sbjct: 171 GLKLGVPELTWSCYEGGDRPCLSCGTCLERTEAFLANGVKDPL 214

Query= sid|114855|lan|dp1ORF034 Phage dp1 ORF|131-652|2
 (173 letters)

>emb|CAB13248| (Z99111) similar to hypothetical proteins [Bacillus subtilis]
 Length = 165

Score = 220 bits (556), Expect = 4e-57
 Identities = 103/139 (74%), Positives = 117/139 (84%)

Query: 5 TTRTDAELTGVTLLGNQDTKYDYDYNPDVLETFPNKHPENNYLVTFDGYEFTSLCPKTGQ 64
 TTR ++EL GVTLGNQ T Y ++Y PDVLE+FPNKH +Y V F+ EFTSLCPKTGQ
 Sbjct: 2 TTRKESELEGVTLLGNQGTNYLFEYAPDVLESFPNKHVNRDYFVKFNCPEFTSLCPKTGQ 61

Query: 65 PDFANVPISYIPNEKMVESKSLKLYLFSFRNHGDFHEDCMNIILNDLYELMEPKYIEVMG 124
 PDFA ++ISYIP+EKMVESKSLKLYLFSFRNHGDFHEDCMNII+NDL ELM+P+YIEV G
 Sbjct: 62 PDFATIYISYIPDEKMVESKSLKLYLFSFRNHGDFHEDCMNIIMNDLIELMDPRYIEVWG 121

Query: 125 LFTPRGGISYIPFVNKVNP 143
 FTPRGGISI P+ N P
 Sbjct: 122 KFTPRGGISIDFYTNYGKP 140

Query= sid|114857|lan|dp1ORF036 Phage dp1 ORF|48808-49362|1
 (184 letters)

>gi|1353529 (U38906) ORF12 [Bacteriophage rlt]
 Length = 296

Score = 53.5 bits (126), Expect = 1e-06
 Identities = 42/149 (28%), Positives = 70/149 (46%), Gaps = 9/149 (6%)

Query: 34 IASNTVNGKTSWAVRLLQRYLAETALDGRIVEKGMFVVSAQLLTFGDYNYFQTMQEFL 93
 + S G GK+ A+ +L+ L T L ++ V + F + + F + + F +
 Sbjct: 155 VVSGPAGTGKSHLAMSILKDCLOHTDLT--VIFASWSEVLHLIKDSFDNKDSFYSTEYFM 212

408

Query: 94 ERFERLKTCELLVIDEIGGSLTKASYPYLYDLVNYRVNNLSTIYTTNYTDDDEIIDLLG 153
 E F + +LLVID+IG +T+ S L ++++ R TI TTN DEI
 Sbjct: 213 EVF---RNTDLLVIDDIGSEKITEWSMSLLTEVLDART----KTIITTNLKSDEIRKKYH 265

Query: 154 QRLYSRIYDTSVVLDQFQASNVRGLEVSEI 182
 R YSR++ F N++ VS++
 Sbjct: 266 NRTYSRLFRGIGKKAPNFENIKDKRVSQ 294

Query= sid|114859|lan|dp1ORF038 Phage dp1 ORF|1350-1871|3
 (173 letters)

>sp|P44123|YB90_HAEIN_HYPOTHETICAL_PROTEIN_H1190 >gi|1074675|pir||F64021 hypothetical
 protein H1190 - Haemophilus influenzae (strain Rd KW20)
 >gi|1574117 (U32798) 6-pyruvoyl tetrahydrobiopterin
 synthase, putative [Haemophilus influenzae Rd]
 Length = 141

Score = 100 bits (247), Expect = 6e-21
 Identities = 59/143 (41%), Positives = 83/143 (57%), Gaps = 10/143 (6%)

Query: 2 RVSKTLTFDAAHQLVGHFGKCANLHGHTYKVEISLAGGTYDHGSSQGMVVDYHVKKIA- 60
 ++SK +FD AH L GH GKC NLHGHTYK+++ ++G Y G+ + MV+DF +K I
 Sbjct: 3 KISKEFSFDMAHLLDGHGDKCQNLHGHTYKQLQVEISGDLYKSGAKKAMVIDFSDLKSIVK 62

Query: 61 GTFIDRLDHAVLL-QGNP-----IALNAVDTKRVLFGRFRTAENMSRFLTWTLTBLMWK 115
 +D +DHA + Q NE L +++K FRITAE ++RF+ L +
 Sbjct: 63 KVILDPMHAFIYDQTNERESQIATLLQKLSKTFGVFPFRTAEIARFIFNRLKH--DE 120

Query: 116 HARIDSIKLWETPTGCAECTYYE 138
 I SI+LWETPT + C Y E
 Sbjct: 121 QLSISSIRLWETPT--SFCEYQE 141

Query= sid|114860|lan|dp1ORF039 Phage dp1 ORF|3306-3803|3
 (165 letters)

>emb|CAA68244| (X99978) ORF7; hydrophobic protein [Lactobacillus plantarum]
 Length = 168

Score = 64.4 bits (154), Expect = 5e-10
 Identities = 49/156 (31%), Positives = 84/156 (53%), Gaps = 9/156 (5%)

Query: 8 WLVRTALIAALYVTLTVAFSAISY--GPIQFRVSEALILLPLWNHRWTFGIVLGTIIANF 65
 W++ AL+AA+YV L + +A S G IQFRVSE L L ++N ++ GIV G I+ +
 Sbjct: 9 WIIN-ALVAMMYVVLCLGPAAFSLASGAIQFRVSEGLNHLAVFNRYIWIWIVAGVILFDA 67

Query: 66 FSP-LGLIDVLFGLSLATFLGXXXXXXXXXXXXSPYSLICPVLA---NAYLIALELRIVY 120
 F P L++VLFG + L ++ + +A + ++IAL + ++
 Sbjct: 68 FCGASLLNVLFGGQSLALLVLTWLPKLTVMQRMLLNIAFTVSMFMIALMITMS 127

Query: 121 S-LPFWESVIYVGISEAIIVLISYFLISTLAKNNHF 155
 S + FW + + +SE II+ I+ ++ +L + HF
 Sbjct: 128 SGVAFWPTYLTALSELIIMSITAPIMYSLDRVLHF 163

Query= sid|114862|lan|dp1ORF041 Phage dp1 ORF|8208-8699|3
 (163 letters)

>gi|2522313 (AF012906) dUTPase homolog [Bacillus subtilis]
 >gi|2634394|emb|CAB13893| (Z99114) similar to
 deoxyuridine 5'-triphosphate nucleotidohydrolase
 [Bacillus subtilis] >gi|3025643 (AF020713) putative
 dUTPase [Bacteriophage SPBc2]
 Length = 142

Score = 108 bits (267), Expect = 2e-23
 Identities = 65/160 (40%), Positives = 83/160 (51%), Gaps = 25/160 (15%)

Query: 5 VDVKMIDPKLDRLLKYT--GDWVDVRISSITKIDADSADVSRCKRVLQKAQVYSVAAGECI 62
 + +K +D R+ GDW+D+R + I D +
 Sbjct: 3 IKIKYLDDETQTRINKMEQGDWIDLRAEDVAIKKDEFKL----- 41

Query: 63 KIAHGFALELPKGYEAILHPRSSLFKKTGLFVSS-GVIDEGYKGDTEWFSVWYATRDA 121
 + G A+ELP+GYEA + PRSS +K G+I +S GVIDE YKGD D WF YA RD
 Sbjct: 42 -VPLGVAMELPEGYEAHVVPSSSTYKNFGVIQTNMGVIDESYKGDNDFFFPAYALRDT 100

409

Query: 122 DIFYDQRIAQFRIQEKQPAIKFNFVESLGNAARGGHGSGTG 161
 I RI QFRI +K PA+ V+ LGN RGGHSGTG
 Sbjct: 101 KIKKGDRIQCFRIMKKMPAVDLIEVDRLGNGDRGGHSGTG 140

Query= sid|114867|lan|dp1ORF046 Phage dp1 ORF|42774-43202|3
 (142 letters)

>emb|CAB07984| (Z93946) hypothetical protein (bacteriophage Dp-1)
 Length = 142

Score = 287 bits (728), Expect = 2e-77
 Identities = 142/142 (100%), Positives = 142/142 (100%)

Query: 1 MPMWLNDTAVLTITACSGVLTVLLNKLFEWKSNAKSVLEDISTTLSTLKQQVDGIDQ 60
 MPMWLNDTAVLTITACSGVLTVLLNKLFEWKSNAKSVLEDISTTLSTLKQQVDGIDQ
 Sbjct: 1 MPMWLNDTAVLTITACSGVLTVLLNKLFEWKSNAKSVLEDISTTLSTLKQQVDGIDQ 60

Query: 61 TTVAINHQNVDVIQDGTQRIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120
 TTVAINHQNVDVIQDGTQRIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE
 Sbjct: 61 TTVAINHQNVDVIQDGTQRIQRYRLYHDLKREVITGYTTLDHFRELSILFESYKNLGGNGE 120

Query: 121 VEALYEKYKKLPIREEDLDETI 142
 VEALYEKYKKLPIREEDLDETI
 Sbjct: 121 VEALYEKYKKLPIREEDLDETI 142

Query= sid|114901|lan|dp1ORF080 Phage dp1 ORF|42490-42759|1
 (89 letters)

>emb|CAB07983| (Z93946) hypothetical protein (bacteriophage Dp-1)
 Length = 124

Score = 147 bits (367), Expect = 1e-35
 Identities = 75/75 (100%), Positives = 75/75 (100%)

Query: 1 MLNLTCSRQIVAEFTIGQGAEEKLVKTTIVNIDANAVSTVSETLHDPDLAANRRELRA 60
 MLNLTCSRQIVAEFTIGQGAEEKLVKTTIVNIDANAVSTVSETLHDPDLAANRRELRA
 Sbjct: 1 MLNLTCSRQIVAEFTIGQGAEEKLVKTTIVNIDANAVSTVSETLHDPDLAANRRELRA 60

Query: 61 EQKLRETRYAIEDEI 75
 EQKLRETRYAIEDEI
 Sbjct: 61 EQKLRETRYAIEDEI 75

Query= sid|114912|lan|dp1ORF091 Phage dp1 ORF|43189-43413|1
 (74 letters)

>emb|CAB07985| (Z93946) holin (bacteriophage Dp-1)
 Length = 74

Score = 63.2 bits (151), Expect = 2e-10
 Identities = 34/74 (45%), Positives = 34/74 (45%)

Query: 1 MKLSNEQYDXXXXXXXXXXXXXXXXXXXXXQFDXXXXXXXXXXXXXXXXXVLGVSSR 60
 MKLSNEQYD YQFD VLGVSSR
 Sbjct: 1 MKLSNEQYDVAKNVVTVVPAALITGLGALYQFDITAITGTIALLATFAGTVLGVS 60

Query: 61 NYQKEQEAQNNEVE 74
 NYQKEQEAQNNEVE
 Sbjct: 61 NYQKEQEAQNNEVE 74

Condensed listing of homology information from above

Phage: dpl

Database: nr

Program: Blastp

Query= sid|114822|lan|dp1ORF001 Phage dpl ORF|36698-40390|2
(1230 letters)

gi 2444124 (U88974) ORF45 [Streptococcus thermophilus temperate ...	467	e-130
gi 928828 (L44593) ORF1904; putative [Lactococcus lactis phage B...	427	e-118
gi 2935676 (AF032121) unknown [Streptococcus thermophilus bacter...	309	1e-82
gi 2935691 (AF032122) unknown [Streptococcus thermophilus bacter...	306	7e-82
gi 3540289 (AF057033) putative anti-receptor [Streptococcus ther...	279	6e-74
gi 4530154 gb AAD21894.1 (AF085222) putative tail-host specific...	220	3e-56
gi 930045 emb CAA33387 (X15332) alpha-1 (III) collagen [Homo sa...	58	4e-07
gi 1070603 pir CGHU7L collagen alpha 1(III) chain precursor - h...	58	4e-07
gi 4502951 ref NP_000081.1 PCOL3A1 collagen, type III, alpha 1 ...	58	4e-07
gi 115290 sp P04258 CA13_BOVIN COLLAGEN ALPHA 1(III) CHAIN >gi 7...	58	4e-07
gi 575322 emb CAA36279 (X52046) type III collagen [Mus musculus]	57	8e-07
gi 2119163 pir S59856 collagen alpha 1(III) chain precursor - m...	57	8e-07
gi 543912 sp P13941 CA13_RAT COLLAGEN ALPHA 1(III) CHAIN >gi 543...	57	1e-06
gi 3171998 emb CAA06510 (AJ005395) collagen alpha 1 (III) [Ratt...	57	1e-06
gi 3947565 emb CAA90250 (Z49967) similar to collagen; cDNA EST ...	54	7e-06
gi 423403 pir A46053 bullous pemphigoid antigen, BPAG2, type XV...	53	9e-06
gi 115410 sp P12114 CCS1_CAEEL CUTICLE COLLAGEN SQT-1 >gi 84437 ...	53	9e-06
gi 3873801 emb CAA90084 (Z49907) cuticle collagen SQT-1; cDNA E...	53	9e-06

Query= sid|114823|lan|dp1ORF002 Phage dpl ORF|32386-35835|1
(1149 letters)

gi 3341922 dbj BAA31888 (AB009866) orf 15 [bacteriophage phi PVL]	280	3e-74
gi 4126622 dbj BAA36642.1 (AB016282) ORF36 [bacteriophage phi-105]	232	1e-59
gi 1369948 emb CAA59194 (X84706) host interacting protein [Bact...	201	3e-50
gi 3139112 (AF063097) gpT [Bacteriophage P2]	188	2e-46
gi 3337272 (U32222) G protein [Bacteriophage 186]	161	3e-38
gi 4063799 dbj BAA36253 (AB008550) orf25; similar to T gene of ...	159	8e-38
gi 3172274 (AF022214) minor tail subunit; putative tape-measure ...	123	6e-27
gi 465127 sp Q05233 VG26_BPML5 MINOR TAIL PROTEIN GP26 >gi 41904...	108	2e-22
gi 3540284 (AF057033) putative minor tail protein [Streptococcus...	99	2e-19
gi 2444119 (U88974) ORF40 [Streptococcus thermophilus temperate ...	90	6e-17
gi 2634555 emb CAB14053 (Z99115) yomI [Bacillus subtilis] >gi 3...	66	1e-09
gi 2392838 (AF011378) unknown [Bacteriophage sk1]	64	5e-09
gi 2764873 emb CAA66557 (X97918) gene 18.1 [Bacteriophage SPP1]	62	3e-08
gi 1353559 (U38906) ORF42 [Bacteriophage rlt]	61	6e-08
gi 630841 pir S39079 puff C-8 protein - fungus gnat (Rhynchosci...	55	2e-06
gi 1730865 sp P51731 YO27_BPHP1 HYPOTHETICAL 72.8 KD PROTEIN IN ...	53	8e-06
gi 224288 prf 1101273J ORF 7 [Bacteriophage HP1]	53	1e-05

Query= sid|114824|lan|dp1ORF003 Phage dpl ORF|53538-55877|3
(779 letters)

gi 118825 sp P00582 DPO1_ECOLI DNA POLYMERASE I (POL I) >gi 6705...	193	3e-48
gi 2982102 pdb 1KFS A Chain A, All-Oxygen Dna Complexed To The 3...	193	3e-48
gi 229889 pdb 1DPI DNA Polymerase I (Klenow Fragment) (E.C.2....	193	3e-48
gi 1169402 sp P43741 DPO1_HAEIN DNA POLYMERASE I (POL I) >gi 107...	191	1e-47
gi 2688462 (AE001156) DNA polymerase I (polA) [Borrelia burgdorf...	190	3e-47
gi 809180 pdb 1KLN A Escherichia coli	190	3e-47
gi 1913934 emb CAA72997 (Y12328) DNA-directed DNA polymerase I ...	189	8e-47
gi 4090935 (AF028719) DNA polymerase type I [Rhodothermus sp. 'I...	175	1e-42
gi 4731571 gb AAD28505.1 AF121780.1 (AF121780) DNA polymerase I ...	174	2e-42
gi 1633576 (U57757) similar to proofreading 3'-5' exonuclease an...	173	4e-42
gi 3322368 (AE001195) DNA polymerase I (polA) [Treponema pallidum]	172	9e-42
gi 1006595 dbj BAA10748 (D64005) DNA polymerase I [Synechocysti...	171	2e-41
gi 585062 sp Q07700 DPO1_MYCTU DNA POLYMERASE I (POL I) >gi 4161...	163	5e-39
gi 4376908 gb AAD18751 (AE001645) DNA Polymerase I [Chlamydia p...	157	2e-37
gi 1169403 sp P46835 DPO1_MYCLE DNA POLYMERASE I (POL I) >gi 107...	152	7e-36
gi 2145839 pir S72949 DNA polymerase I - Mycobacterium leprae >...	152	7e-36
gi 1405438 emb CAA67184 (X98575) DNA-dependent DNA polymerase [...	152	9e-36
gi 2506365 sp P80194 DPO1_THECA DNA POLYMERASE I, THERMOSTABLE (...	147	2e-34
gi 3328929 (AE001322) DNA Polymerase I [Chlamydia trachomatis]	147	3e-34

411

gi 3913510 sp O52225 DPO1_THEFI DNA POLYMERASE I, THERMOSTABLE (...)	146	7e-34
gi 1205984 (U33536) DNA polymerase I [Bacillus stearothermophilus]	146	7e-34
gi 118827 sp P13252 DPO1_STRPN DNA POLYMERASE I (POL I) >gi 9802...	145	9e-34
gi 1942202 pdb 1JXE Stoffel Fragment Of Taq Dna Polymerase I	145	1e-33
gi 1943520 pdb 1KTQ Dna Polymerase	145	1e-33
gi 1084022 pir JX0359 DNA-directed DNA polymerase (EC 2.7.7.7) ...	145	1e-33
gi 507891 dbj BAA06775 (D32013) DNA Polymerase [Thermus aquaticus]	145	1e-33
gi 118828 sp P19821 DPO1_THEAQ DNA POLYMERASE I, THERMOSTABLE (T...	145	1e-33
gi 1706502 sp P52028 DPO1_THETH DNA POLYMERASE I, THERMOSTABLE (...)	144	2e-33
gi 1097211 prf 2113329A DNA polymerase [Thermus aquaticus therm...	144	2e-33
gi 2098289 pdb 1TAU A Chain A, Structure Of Dna Polymerase	143	3e-33

Query= sid|114825|lan|dp1ORF004 Phage dp1 ORF|40401-42440|3
(679 letters)

gi 1934761 emb CAB07981 (Z93946) hypothetical protein [bacterio...	1011	0.0
gi 3540290 (AF057033) putative minor structural protein [Strepto...	346	2e-94
gi 2444125 (U88974) ORF46 [Streptococcus thermophilus temperate ...]	339	3e-92
gi 1934762 emb CAB07982 (Z93946) hypothetical protein [bacterio...	300	2e-80
gi 4530155 gb AAD21895.1 (AF085222) unknown [Streptococcus ther...	276	4e-73
gi 2935677 (AF032121) unknown [Streptococcus thermophilus bacter...	250	3e-65
gi 2935692 (AF032122) unknown [Streptococcus thermophilus bacter...	250	3e-65
gi 1136289 (U42597) histidine kinase A [Dictyostelium discoideum]	50	7e-05

Query= sid|114827|lan|dp1ORF006 Phage dp1 ORF|45296-46987|2
(563 letters)

gi 4377165 gb AAD18987 (AE001666) SWI/SNF family helicase_2 [Ch...	171	1e-41
gi 1769947 emb CAA67095 (X98455) SNF [Bacillus cereus]	160	3e-38
gi 3329163 (AE001341) SWF/SNF family helicase [Chlamydia trachom...	159	6e-38
gi 4377149 gb AAD18973 (AE001664) SWI/SNF family helicase_1 [Ch...	157	2e-37
gi 3328995 (AE001326) SWI/SNF family helicase [Chlamydia trachom...	153	2e-36
gi 2493354 sp P75093 Y018_MYCPN HYPOTHETICAL HELICASE MG018/MG01...	146	4e-34
gi 1653748 dbj BAA18659 (D90916) helicase of the snf2/rad54 fam...	143	3e-33
gi 1763712 emb CAB05939 (Z83337) member of the SNF2 helicase fa...	143	4e-33
gi 2636153 emb CAB15645.1 (Z99122) similar to SNF2 helicase [Ba...	143	4e-33
gi 2909552 emb CAA17284 (AL021924) helZ [Mycobacterium tubercul...	140	2e-32
gi 3844627 (U39681) ATP-dependent RNA helicase, putative [Mycopla...	136	3e-31
gi 1351463 sp P47264 Y018_MYCGE HYPOTHETICAL HELICASE MG018	136	4e-31
gi 2660669 (AC002342) human Mi-2 autoantigen-like protein [Arabi...	131	2e-29
gi 1361537 pir I64201 helicase (mot1) homolog - Mycoplasma geni...	129	4e-29
gi 3482977 emb CAA20533.1 (AL031369) putative protein [Arabidop...	128	9e-29
gi 3298562 (U91543) zinc-finger helicase [Homo sapiens]	120	2e-26
gi 3875971 emb CAB02491 (Z80344) similar to helicase; cDNA EST ...	120	2e-26
gi 4557451 ref NP_001263.1 PCHD3 chromodomain helicase DNA bind...	120	2e-26
gi 2645435 (AF007780) CHD3 [Drosophila melanogaster]	118	1e-25
gi 3875165 emb CAA91798 (Z67881) Similarity to Mouse Chromodoma...	118	1e-25

Query= sid|114828|lan|dp1ORF007 Phage dp1 ORF|22230-23621|3
(463 letters)

gi 2444105 (U88974) ORF26 [Streptococcus thermophilus temperate ...]	89	7e-17
gi 3318666 (U19754) BBA31 homolog [Borrelia burgdorferi]	59	7e-08
gi 2690260 (AE000790) conserved hypothetical protein [Borrelia b...	56	5e-07

Query= sid|114829|lan|dp1ORF008 Phage dp1 ORF|49624-50961|1
(445 letters)

gi 4406210 gb AAD19901 (AF100420) DnaB replication fork helicase...	68	2e-10
gi 3121983 sp O25916 DNAB_HELPY REPLICATIVE DNA HELICASE >gi 231...	67	2e-10
gi 4416322 gb AAD20314 (AF106032) replicative helicase; DnaB [B...	65	9e-10
gi 4155895 (AE001551) REPLICATIVE DNA HELICASE [Helicobacter pyl...	60	4e-08
gi 3322317 (AE001191) replicative DNA helicase (dnaB) [Treponema...	58	1e-07
gi 1380311 sp P04530 VG41 PRIMASE-HELICASE (PROTEIN GP41) >g...	53	3e-06
gi 2983861 (AE000742) replicative DNA helicase [Aquifex aeolicus]	51	1e-05

Query= sid|114831|lan|dp1ORF010 Phage dp1 ORF|8699-9859|2
(386 letters)

gi 2760912 (AF037258) RecA protein [Chlorobium tepidum]	133	2e-30
gi 3219851 sp P94666 RECA_CLOPE RECA PROTEIN >gi 1698591 (U61497...	129	3e-29
gi 1350566 sp P48295 RECA_STRVL RECA PROTEIN >gi 508860 (U04837)...	128	7e-29
gi 744163 prf 2014250A recA-like protein [Streptomyces violaceus]	126	3e-28
gi 730487 sp P41054 RECA_STRAM RECA PROTEIN >gi 511133 emb CAA82...	125	4e-28
gi 2687334 emb CAA15875 (AL020958) RecA protein [Streptomyces c...	125	6e-28
gi 1350565 sp P48294 RECA_STRLI RECA PROTEIN >gi 481482 pir S38...	125	6e-28

412

gi|464599|sp|P33542|RECA_AQUFY RECA PROTEIN >gi|1086167|pir||A55... 123 2e-27
 gi|417636|sp|P32725|RECA_RHOSH RECA PROTEIN >gi|541307|pir||S415... 123 2e-27
 gi|2984348 (AE000775) recombination protein RecA [Aquifex aeolicus] 123 2e-27
 gi|3219854|sp|P95846|RECA_STRRM RECA PROTEIN >gi|1729800|emb|CAA... 122 4e-27
 gi|2500086|sp|Q59560|RECA_MYCSM RECA PROTEIN >gi|1430892|emb|CAA... 122 4e-27
 gi|1350567|sp|P48296|RECA_THEAQ RECA PROTEIN >gi|1072963|pir||A5... 122 6e-27
 gi|625663|pir||JX0292 recA protein - Thermus aquaticus (strain HB8) 121 1e-26
 gi|1172880|sp|P42440|RECA_CAMJE RECA PROTEIN >gi|2119991|pir||I4... 120 2e-26
 gi|4154654 (AE001453) RECA PROTEIN. [Helicobacter pylori J99] 120 2e-26
 gi|1072968|pir||C55020 recA protein - Thermus sp >gi|458472|dbj|... 120 2e-26
 gi|3219852|sp|P95469|RECA_PARDE RECA PROTEIN >gi|1825468 (U59631... 119 3e-26
 gi|2507284|sp|P42445|RECA_HELPY RECA PROTEIN >gi|2313235|gb|AAD0... 119 4e-26
 gi|1172890|sp|Q02350|RECA_STAAU RECA PROTEIN >gi|463285 (L25893)... 118 5e-26
 gi|4416209|gb|AAD20261| (AF094756) RecA protein [Bifidobacterium... 118 5e-26
 gi|2500084|sp|Q59180|RECA_BORBU RECA PROTEIN >gi|1276443 (U23457... 118 5e-26

Query= sid|114832|lan|dp1ORF011 Phage dpl ORF|28017-29096|3
 (359 letters)

gi|2444110 (U88974) ORF31 [Streptococcus thermophilus temperate ... 187 1e-46
 gi|3320438 (AF057033) gp348 [Streptococcus thermophilus bacterio... 179 2e-44
 gi|479514|pir||S34244 hypothetical protein p38 - actinophage VWB... 62 8e-09

Query= sid|114834|lan|dp1ORF013 Phage dpl ORF|10215-11240|3
 (341 letters)

gi|580855|emb|CAA29958| (X06803) dnaZX-like ORF put. DNA polymer... 182 2e-45
 gi|118807|sp|P09122|DP3X_BACSU DNA POLYMERASE III SUBUNITS GAMMA... 182 2e-45
 gi|98292|pir||S13786 DNA-directed DNA polymerase (EC 2.7.7.7) II... 182 2e-45
 gi|1527142 (U66040) DNA polymerase III gamma subunit [Salmonella... 172 4e-42
 gi|2494197|sp|P74876|DP3X_SALTY DNA POLYMERASE III SUBUNITS GAMM... 172 4e-42
 gi|118808|sp|P06710|DP3X_ECOLI DNA POLYMERASE III SUBUNITS GAMMA... 170 1e-41
 gi|4155207 (AE001497) DNA POLYMERASE III SUBUNITS GAMMA AND TAU ... 169 2e-41
 gi|2313841|gb|AAD07767.1| (AE000584) DNA polymerase III gamma an... 168 4e-41
 gi|2583049 (AF025391) DNA polymerase III holoenzyme tau subunit ... 166 3e-40
 gi|2984127 (AE000759) DNA polymerase III gamma subunit [Aquifex ... 166 3e-40
 gi|3861390|emb|CAA15289| (AJ235273) DNA POLYMERASE III SUBUNITS ... 165 5e-40
 gi|1169397|sp|P43746|DP3X_HAEIN DNA POLYMERASE III SUBUNITS GAMM... 156 2e-37
 gi|1293572 (U49738) DNA polymerase III tau homolog DnaX [Cauloba... 151 8e-36
 gi|3328753 (AE001306) DNA Pol III Gamma and Tau [Chlamydia trach... 148 4e-35
 gi|4376294|gb|AAD18193| (AE001589) DNA Polymerase III Gamma and ... 148 5e-35
 gi|581255|emb|CAA28175| (X04487) alternate dnaZX protein (AA 1-6... 146 3e-34
 gi|2688379 (AE001151) DNA polymerase III, subunits gamma and tau... 140 2e-32
 gi|3323329 (AE001268) DNA polymerase III, subunits gamma and tau... 137 1e-31

Query= sid|114835|lan|dp1ORF014 Phage dpl ORF|50961-51974|3
 (337 letters)

gi|1346796|sp|P47492|PRIM_MYCGE DNA PRIMASE >gi|1361496|pir||F64... 57 2e-07
 gi|740008|prf||2004290A primase [Haemophilus influenzae] 51 1e-05
 gi|1172619|sp|Q08346|PRIM_HAEIN DNA PRIMASE >gi|1074033|pir||A64... 51 1e-05
 gi|1709769|sp|Q04505|PRIM_LACLA DNA PRIMASE >gi|1075726|pir||JC2... 51 1e-05
 gi|639846|dbj|BAA03516| (D14690) DNA primase [Lactococcus lactis] 51 1e-05

Query= sid|114837|lan|dp1ORF016 Phage dpl ORF|43413-44303|3
 (296 letters)

gi|1934766|emb|CAB07986| (Z93946) N-acetylmuramoyl-L-alanine ami... 661 0.0
 gi|113676|sp|P06653|ALYS_STRPN AUTOLYSIN (N-ACETYLMURAMOYL-L-ALA... 221 4e-57
 gi|282326|pir||A42935 N-acetylmuramoyl-L-alanine amidase (EC 3.5... 219 3e-56
 gi|416618|sp|P32762|ALYS_BPHB3 LYTC AMIDASE (N-ACETYLMURAMOYL-L... 212 2e-54
 gi|285273|pir||A42936 N-acetylmuramoyl-L-alanine amidase (EC 3.5... 212 2e-54
 gi|127787|sp|P15057|LYCA_BPCP1 LYSOZYME (ENDOLYSIN) (MURAMIDASE)... 162 4e-39
 gi|67761|pir||MUBPCP N-acetylmuramoyl-L-alanine amidase (EC 3.5... 162 4e-39
 gi|127789|sp|P19386|LYCA_BPCP9 LYSOZYME (ENDOLYSIN) (MURAMIDASE)... 160 1e-38
 gi|928832 (L44593) ORF259; putative [Lactococcus lactis phage BK... 119 2e-26
 gi|2511705|emb|CAA71783| (Y10818) sigA binding protein [Streptoc... 111 9e-24
 gi|4097980 (U72655) surface protein C [Streptococcus pneumoniae] 107 1e-22
 gi|2351768 (U89711) PspA [Streptococcus pneumoniae] 105 4e-22
 gi|2425109 (AF019904) choline binding protein A [Streptococcus p... 104 6e-22
 gi|282335|pir||A41971 surface protein pspA precursor - Streptoco... 104 1e-21
 gi|2576331|emb|CAA05158| (AJ002054) SpsA protein [Streptococcus ... 103 2e-21
 gi|2127295|pir||S57962 cspC protein - Clostridium acetobutylicum... 85 6e-16
 gi|2576333|emb|CAA05159| (AJ002055) SpsA protein [Streptococcus ... 84 1e-15
 gi|4106522|gb|AAD02874.1| (AF097909) excreted protein FibB [Pept... 83 3e-15
 gi|1361406|pir||S57714 cspB protein - Clostridium acetobutylicum... 82 4e-15
 gi|1914872|emb|CAB04758| (Z82001) PCPA [Streptococcus pneumoniae] 81 9e-15

gi 3168594 dbj BAA28613 (AB012763) SpaA (Erysipelothrix rhusiop...	81	1e-14
gi 2292750 emb CAA64942 (X95646) homology to orf259 of lactococ...	80	3e-14
gi 2935696 (AF032122) putative lysin [Streptococcus thermophilus...	80	3e-14
gi 4586910 dbj BAA76540.1 (AB017447) protective antigen SpaA.1 ...	80	3e-14
gi 3540294 (AF057033) lysin [Streptococcus thermophilus bacterio...	79	5e-14

Query= sid|114841|lan|dp1ORF020 Phage dp1 ORF|1864-2658|1
(264 letters)

gi 2633745 emb CAB13247 (Z99111) similar to coenzyme PQQ synthe...	217	5e-56
gi 2808502 emb CAA12532 (AJ225561) ExsD protein [Sinorhizobium ...	163	1e-39
gi 3861151 emb CAA15051 (AJ235272) unknown [Rickettsia prowazekii]	82	6e-15
gi 1652793 dbj BAA17712 (D90908) hypothetical protein [Synechoc...	76	3e-13
gi 1723815 sp P55139 YGCF_ECOLI HYPOTHETICAL 25.0 KD PROTEIN IN ...	70	2e-11
gi 2984272 (AE000769) hypothetical protein [Aquifex aeolicus]	66	4e-10
gi 4155435 (AE001516) putative [Helicobacter pylori J99]	57	1e-07
gi 2127833 pir C64505 coenzyme PQQ synthesis protein III homolo...	55	5e-07
gi 2622338 (AE000890) coenzyme PQQ synthesis protein III [Methan...	54	9e-07
gi 3257042 dbj BAA29725 (AP000003) 254aa long hypothetical prot...	53	2e-06
gi 2314068 gb AAD07976.1 (AE000602) conserved hypothetical prot...	52	6e-06
gi 1723816 sp P45097 YGCF_HAEIN HYPOTHETICAL PROTEIN H1189 >gi ...	50	2e-05

Query= sid|114842|lan|dp1ORF021 Phage dp1 ORF|2504-3295|2
(263 letters)

gi 127481 sp P19465 GCH1_BACSU GTP CYCLOHYDROLASE I (GTP-CH-I) >...	208	4e-53
gi 3242315 emb CAA04237 (AJ000685) GTP cyclohydrolase [Streptoc...	191	4e-48
gi 2494695 sp Q54769 GCH1_SYN7 GTP CYCLOHYDROLASE I (GTP-CH-I) ...	189	2e-47
gi 255061 bbs 112832 (S44049) GTP cyclohydrolase I (clone hGCH-1...	187	7e-47
gi 4503949 ref NP_000152.1 PGCH1 GTP cyclohydrolase 1 (dopa-res...	187	7e-47
gi 2113967 emb CAB08935 (Z95557) fole [Mycobacterium tuberculosis]	187	7e-47
gi 1730240 sp P50141 GCH1_CHICK GTP CYCLOHYDROLASE I (GTP-CH-I) ...	185	3e-46
gi 2494696 sp Q55759 GCH1_SYN3 GTP CYCLOHYDROLASE I (GTP-CH-I) ...	184	5e-46
gi 121061 sp P22288 GCH1_RAT GTP CYCLOHYDROLASE I PRECURSOR (GTP...	184	6e-46
gi 3183014 sp O13774 GCH1_SCHPO GTP CYCLOHYDROLASE I (GTP-CH-I) ...	184	6e-46
gi 3097224 emb CAA18795 (AL023093) GTP cyclohydrolase I [Mycoba...	182	2e-45
gi 2494697 sp Q19980 GCH1_CAEEL PROBABLE GTP CYCLOHYDROLASE I (G...	182	2e-45
gi 462167 sp Q05915 GCH1_MOUSE GTP CYCLOHYDROLASE I PRECURSOR (G...	180	7e-45
gi 1669664 emb CAA89808 (Z49706) GTP cyclohydrolase I [Dictyost...	180	1e-44
gi 2981082 (AF052048) GTP-cyclohydrolase [Ostertagia ostertagi]	178	3e-44
gi 31954 emb CAA78908 (Z16418) GTP cyclohydrolase I [Homo sapi...	177	8e-44
gi 551344 bbs 150280 (S71373) GTP cyclohydrolase I [mice, Peptid...	174	5e-43
gi 1730247 sp P51601 GCH1_YEAST GTP CYCLOHYDROLASE I (GTP-CH-I) ...	174	7e-43
gi 1246912 emb CAA87397 (Z47201) GTP cyclohydrolase 1 [Saccharo...	172	2e-42
gi 1730246 sp P51595 GCH1_STRPN GTP CYCLOHYDROLASE I (GTP-CH-I) ...	168	3e-41
gi 2982951 (AE000680) GTP cyclohydrolase I [Aquifex aeolicus]	164	6e-40

Query= sid|114843|lan|dp1ORF022 Phage dp1 ORF|30896-31675|2
(259 letters)

gi 2347102 (U77367) internalin [Listeria monocytogenes]	55	5e-07
gi 3123226 sp P25146 INLA_LISMO INTERNALIN A PRECURSOR >gi 48705...	52	4e-06
gi 149674 (M67471) internalin [Listeria monocytogenes]	52	4e-06

Query= sid|114850|lan|dp1ORF029 Phage dp1 ORF|662-1348|2
(228 letters)

gi 2650185 (AE001074) succinoglycan biosynthesis regulator (exsB...	119	2e-26
gi 3861231 emb CAA15131 (AJ235272) unknown [Rickettsia prowazekii]	117	8e-26
gi 2622210 (AE000881) conserved protein [Methanobacterium thermo...	108	4e-23
gi 2983380 (AE000709) trans-regulatory protein ExsB [Aquifex aeo...	88	6e-17
gi 1001327 dbj BAA10814 (D64006) ExsB [Synechocystis sp.]	88	6e-17
gi 2128055 pir B64468 hypothetical protein homolog MJ1347 - Met...	83	1e-15
gi 4155143 (AE001491) putative [Helicobacter pylori J99]	82	4e-15
gi 2313760 gb AAD07701.1 (AE000578) conserved hypothetical prot...	80	2e-14
gi 2120814 pir S60183 protein ExsB - Rhizobium meliloti >gi 114...	76	3e-13
gi 2633743 emb CAB13245 (Z99111) similar to hypothetical protei...	75	5e-13
gi 1175543 sp P44124 YBAX_HAEIN HYPOTHETICAL PROTEIN H11191 >gi ...	74	1e-12
gi 2495537 sp P77756 YBAX_ECOLI HYPOTHETICAL 25.5 KD PROTEIN IN ...	71	5e-12
gi 3256471 dbj BAA29154.1 (AP000001) 269aa long hypothetical pr...	67	1e-10
gi 2921156 (AF022216) aluminum resistance protein [Arthrobacter ...	54	1e-06

Query= sid|114855|lan|dp1ORF034 Phage dp1 ORF|131-652|2
(173 letters)

gi 2633746 emb CAB13248 (Z99111) similar to hypothetical protei...	220	4e-57
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-11-

gi|4155926 (AE001554) putative [Helicobacter pylori J99] 162 1e-39
gi|2314588|gb|AAD08456.1| (AE000642) conserved hypothetical prot... 161 3e-39
gi|2983458 (AE000714) hypothetical protein [Aquifex aeolicus] 103 9e-22
gi|1006604|dbj|BAA10757| (D64005) hypothetical protein [Synechoc... 87 6e-17
gi|2967529 (U11045) unknown [Buchnera aphidicola] 79 2e-14
gi|2495654|sp|Q46920|YQCD_ECOLI HYPOTHETICAL 32.6 KD PROTEIN IN ... 69 2e-11
gi|1175604|sp|P44153|YQCD_HAEIN HYPOTHETICAL PROTEIN HI1291 >gi|... 63 1e-09
gi|3860642|emb|CAA14543| (AJ235270) unknown [Rickettsia prowazekii] 56 1e-07

Query= sid|114857|lan|dp1ORF036 Phage dp1 ORF|48808-49362|1
(184 letters)

gi|1353529 (U38906) ORF12 [Bacteriophage rlt] 53 1e-06

Query= sid|114859|lan|dp1ORF038 Phage dp1 ORF|1350-1871|3
(173 letters)

gi|1175542|sp|P44123|YB90_HAEIN HYPOTHETICAL PROTEIN HI1190 >gi|... 100 6e-21
gi|2982977 (AE000681) hypothetical protein [Aquifex aeolicus] 67 7e-11
gi|3860744|emb|CAA14645| (AJ235270) unknown [Rickettsia prowazekii] 65 3e-10
gi|2650193 (AE001074) conserved hypothetical protein [Archaeoglo... 58 4e-08
gi|3258383|dbj|BAA31066.1| (AP000007) 157aa long hypothetical pr... 55 2e-07
gi|1001713|dbj|BAA10550| (D64004) hypothetical protein [Synechoc... 50 8e-06
gi|4155434 (AE001516) putative [Helicobacter pylori J99] 50 1e-05

Query= sid|114860|lan|dp1ORF039 Phage dp1 ORF|3306-3803|3
(165 letters)

gi|1922884|emb|CAA68244| (X99978) ORF7; hydrophobic protein [Lact... 64 5e-10

Query= sid|114862|lan|dp1ORF041 Phage dp1 ORF|8208-8699|3
(163 letters)

gi|2522313 (AF012906) dUTPase homolog [Bacillus subtilis] >gi|26... 108 2e-23
gi|2634150|emb|CAB13650| (Z99113) similar to deoxyuridine 5'-tri... 108 3e-23
gi|3913546|sp|O54134|DUT_STRCO DEOXYURIDINE 5'-TRIPHOSPHATE NUCL... 56 2e-07
gi|3913542|sp|O48500|DUT_BPTS DEOXYURIDINE 5'-TRIPHOSPHATE NUCLE... 52 3e-06
gi|3913548|sp|O68992|DUT_CHLTE DEOXYURIDINE 5'-TRIPHOSPHATE NUCL... 50 1e-05

Query= sid|114867|lan|dp1ORF046 Phage dp1 ORF|42774-43202|3
(142 letters)

gi|1934764|emb|CAB07984| (Z93946) hypothetical protein [bacterio... 287 2e-77

Query= sid|114901|lan|dp1ORF080 Phage dp1 ORF|42490-42759|1
(89 letters)

gi|1934763|emb|CAB07983| (Z93946) hypothetical protein [bacterio... 147 1e-35

Query= sid|114912|lan|dp1ORF091 Phage dp1 ORF|43189-43413|1
(74 letters)

gi|1934765|emb|CAB07985| (Z93946) holin [bacteriophage Dp-1] 63 2e-10

Table 32

Sequence of Dp1 published by Sheehan and al.. 4731 nucleotides.

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141	agaccttaaa	tatcgaattg	actcaaaagc	cgatcaaaag	ctaactaacc	aacagttgac	ggcactcacg
211	gaaaaggctc	aactacatga	cgcagaactg	aaagctaagg	ctacaatgga	gcagtttaagt	aacttagaaa
281	aggcttatga	aggtagaatg	aaagctaagt	aagaagctat	caacaaatcg	gaaccggacc	taatcttagc
351	ggcaagtcca	attgaagcta	ctatccaaga	acttggcggg	ctacgggaac	tgaagaagtt	cgtcgacagt
421	tgcagtagct	cttctaatac	agggtctaatt	atcggtaaga	acgacggtag	ctctaccatt	aagggtatcaa
491	gtgaccgaat	ttctatgttc	tccgcaggga	atgaagttat	gtaccttacg	caagggttca	ttcacatcga
561	taacgggatac	tttaccceat	ccattcaagt	cgggccgattt	agaacgggaac	aatactcggt	taatccagac
631	atgaacgtga	ttcggtatgt	aggataagga	gaataacatg	acaaaattta	tcaactcata	cggccctctt
701	cacttgaacc	tttaccgtcga	acaagtttagt	caggacgtta	cgaacaactc	ctcgcgaggt	agttggcgag
771	ctactgtcga	ccgcgatgga	gcttatcgaa	cgtggactta	tggaaatatt	agtaaccttt	ccgtatgggt
841	aaatgggttca	agtgttcata	gcagtcaccc	agactacgac	acgtccggcg	aagaggttaac	gctcgcaagt
911	ggagaagtga	ctgttcctca	caatagttag	gggacaaaaga	caatgtccgt	ttgggcttcg	tttgacccta
981	ataacggcgt	tcaacggaaat	atcactatct	ctactaatta	cacttttagac	agtattccaa	ggtctacaca
1051	gatttctagt	tttgaggga	atcgaaatct	aggatcttta	catacgggtta	tctttaaccg	aaaagtgaac
1121	tctttttacg	atcaagtttg	gtaccgagtt	ttcggtagcg	actggataga	tttaggttaag	aacctacta
1191	ctagcgtatc	ctttacgcgc	tcaactggact	tagcaaggta	cttacctaaa	tcaagttccg	gaacaatgga
1261	catctgtatt	cgaacctata	acggaaactac	gcaaatgggt	agtgcagctt	attcaaacgg	atggagggtc
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1401	agattttaac	agggaaacaac	ttctccaaa	tcagtgtcgaa	cattcaagtc	aacttcaaca	atgcttcggg
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1821	cagaagatag	aggttcgccg	tcaggagcgt	tcaactactat	ttccctactg	actaactcgt	ccgcgaactt
1891	agctgttaac	tacggggccg	acaagtctta	catagttaag	gctaaaaatcc	aagacaggtt	cacttcgact
1961	gaatttagtg	ctacgggtacc	taccgaatca	gtagtcttta	actatgacaa	ggacggtcga	cttggaggtg
2031	gtaagggtgt	agaacaagg	aaggcagggt	caattgatgc	agcagggtgat	atatatgctg	gagggtcgaca
2101	agttcaacag	tttcagctca	ctgataataa	tggagcattg	aacaggggtc	aatataacga	tgttggataa
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2241	gggactattt	caaaatttct	ggtagatag	ctggaaaaatg	gttcaatcct	tcattacaat	gtcaggaaga
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2381	aagacttcga	acagaataat	tggcagaaac	ttgttcttca	aagtgggttg	aacctcact	caacctatgg
2451	cgacgcattc	tattcgaaaa	ctcttgacgg	catagtatat	ttgagaggaa	atgtgcataa	aggacttatc
2521	gacaaaacg	ctactattgc	agtacttctc	gaaggattta	gaccgaaagt	ttcaatgtat	cttcaggctc
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2661	agataattct	tgggttaaat	tagacaatgt	ctcatttctg	attttaattg	agctgaaatc	atgttataat
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2871	ccgtctctga	aactcttcat	gacccagact	tgtatgctgc	gaaccgtcga	gaacttcgag	ctgacgagca
2941	aaaacttcgc	gaaactcgtt	acgcaatcga	agatgaaatt	aatagctggc	gcggggggaa	aaagggggag
3011	cccggctcta	acaggctgaa	taaggaggcg	tcaatctatg	ccaatgtggc	tacacgacac	cgcagctctg
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3291	taccgtcttt	atcacgactt	aaaaagggaa	gtgataacag	gctatacaac	tctcgaccat	tttagagagc
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3641	gaaactacca	aaaggaacaa	gaagctcaaa	acaatgaggt	ggaataatgg	gagtcgatat	tgaaaaaggc
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Table 33

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gi|153702|gb|J01796.1|STRMALMXP [153702]
gi|153701|gb|J01795.1|STRMALMX [153701]
gi|153693|gb|M13812.1|STRLYTPN [153693]
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gi|398102|gb|L20564.1|STREXP9B [398102]
gi|398100|gb|L20563.1|STREXP9A [398100]
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gi|153626|gb|J04234.1|STREXOA [153626]
gi|153612|gb|M11226.1|STRDPNM [153612]
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gi|153601|gb|M25526.1|STRDN87577 [153601]
gi|153599|gb|M25522.1|STRDN179 [153599]
gi|153594|gb|M37688.1|STRDACA [153594]
gi|153582|gb|L07752.1|STRATTB [153582]
gi|466514|gb|L31413.1|STR1RRA [466514]
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gi|153539|gb|M25516.1|STR110K70 [153539]
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CLAIMS

What is claimed is:

5 1. A method for identifying a bacteriophage coding region encoding a product active on an essential bacterial target, comprising identifying a nucleic acid sequence encoding a gene product which provides a bacteria-inhibiting function when said bacteriophage infects a host bacterium,
 wherein said bacteriophage is uncharacterized and said host bacterium
10 is a pathogenic bacterium.

 2. The method of claim 1, further comprising expressing a recombinant bacteriophage ORF in cells of a bacterial strain, wherein inhibition of said cells following expression of said ORF is indicative that said product is active on an
15 essential bacterial target.

 3. The method of claim 2, wherein inhibition of said bacterium following expression of said ORF is determined by comparison with the growth or viability of said bacterium following expression of an inactivated mutant form of said ORF or in
20 the absence of expression of said ORF, and wherein inhibition of said bacterium following expression of said ORF is indicative that said product is active on an essential bacterial target.

 4. The method of claim 2, wherein expression of said ORF is inducible.
25

 5. The method of claim 1, further comprising sequencing at least a portion of a bacteriophage genome.

 6. The method of claim 1, wherein at least a portion of the nucleotide
30 sequence of a bacteriophage genome is known, said method further comprising identifying at least one ORF in said portion by computer analysis of said sequence.

 7. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify
35 homologous genes or gene products of known biochemical function, thereby indicating the biochemical function of said polypeptide.

8. The method of claim 7, wherein said homologous gene or gene product is a bacterial gene important for cell viability.

9. The method of claim 7, wherein said homologous gene or gene product is a gene or gene product known to have a bacteria-inhibiting function.

10. The method of claim 6, further comprising analyzing the sequence of said at least one ORF or of a polypeptide encoded by said ORF to identify structural motifs in said polypeptide, thereby indicating the cellular function of said polypeptide.

11. The method of claim 1, wherein a host bacterium for said bacteriophage is selected from the species group consisting of bacteria listed in Table 1.

12. The method of claim 1, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

13. The method of claim 2, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.

14. The method of claim 13, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.

15. The method of claim 14, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.

16. The method of claim 1, wherein said pathogenic bacterium is an animal pathogen.

17. The method of claim 16, wherein said pathogenic bacterium is a human pathogen.

18. The method of claim 1, wherein said pathogenic bacterium is a plant pathogen.

19. The method of claim 1, further comprising confirming the inhibitor function of said ORF.

20. The method of claim 19, wherein said confirming comprises expressing a loss-of-function mutant form of said ORF in said host bacterium.

5 21. The method of claim 1, wherein said identifying a nucleic acid sequence encoding a gene product active on an essential bacterial target comprises identifying a nucleic acid sequence encoding a homolog of a bacteriophage polypeptide known to be active on an essential bacterial target.

10 22. The method of claim 1, wherein said identifying a bacteriophage coding region comprises identifying a first coding region from a bacteriophage having a non-pathogenic host bacterial strain related to said pathogenic bacterium, said first coding region encoding a product active on an essential bacterial target; and
 identifying a homolog of said first coding region, wherein said
15 homolog is a probable said bacteriophage coding region encoding a product active on an essential bacterial target.

23. The method of claim 2, wherein a plurality of bacteriophage ORFs from a plurality of different bacteriophage are expressed in at least one bacterium.
20

24. The method of claim 23, wherein each of said plurality of bacteriophage ORFs are expressed in different bacteria.

25 25. A method for identifying a target for antibacterial agents, comprising determining the bacterial target of an uncharacterized bacteriophage inhibitor protein.

26. The method of claim 25, wherein said determining comprises identifying at least one bacterial protein which binds to said bacteriophage inhibitor
30 protein or a fragment thereof.

27. The method of claim 26, wherein said binding is determined using affinity chromatography on a solid matrix.

35 28. The method of claim 25, wherein said determining comprises identifying at least one protein:protein interaction using a genetic screen.

29. The method of claim 28, wherein said genetic screen is a yeast two-hybrid screen.

30. The method of claim 25, wherein said determining comprises a co-immunoprecipitation assay or a protein-protein crosslinking assay.

31. The method of claim 25, wherein said determining comprises identifying a mutated bacterial coding sequence which protects a bacterium from said bacteriophage inhibitor.

32. The method of claim 25, wherein said determining comprises identifying a bacterial coding sequence which protects a bacterium against said bacteriophage inhibitor when expressed at high levels in said bacterium.

33. The method of claim 25, wherein said determining further comprises identifying a bacterial nucleic acid sequence encoding a polypeptide target of said bacteriophage inhibitor protein.

34. The method of claim 33, wherein said nucleic acid sequence is identified by determining at least a portion of the amino acid sequence of a bacterial protein target, and identifying a bacterial nucleic acid sequence which encodes said protein target.

35. The method of claim 25, wherein said bacterial target is naturally produced by a bacterial species selected from the group consisting of species of the genera listed in Table 1.

36. The method of claim 25, wherein said bacterial target is naturally produced by a bacterial strain selected from the group consisting of species listed in Table 1.

37. The method of claim 25, wherein said inhibitor protein is naturally produced by a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

38. The method of claim 25, further comprising identifying a bacteriophage ORF which encodes a product having a bacteria-inhibiting function.

39. The method of claim 38, wherein said identifying a phage ORF comprises expressing at least one bacteriophage ORF in a bacterium, wherein inhibition of said bacterium following said expression is indicative that said ORF
5 encodes a bacteria-inhibiting function.

40. The method of claim 39, wherein a plurality of bacteriophage ORFs are expressed in at least one bacterium.

10 41. The method of claim 40, wherein each of said plurality of bacteriophage ORFs is expressed in a different bacterium.

42. The method of claim 41, wherein said plurality of bacteriophage ORFs comprises at least 10% of the ORFs in the genome of said bacteriophage.
15

43. The method of claim 25, wherein said determining the bacterial target of a bacteriophage inhibitor protein is performed for a plurality of different bacteriophage of the same host bacterium.

20 44. The method of claim 25, wherein said bacterial target originates from an animal pathogen.

45. The method of claim 44, wherein said bacterial target is a gene homologous to a gene from an animal pathogen.
25

46. The method of claim 44, wherein said pathogen is a human pathogen.

47. The method of claim 25, wherein said bacterial target originates from a plant pathogen.
30

48. The method of claim 25, wherein said bacterial target is a gene homologous to a gene from a plant pathogen.

49. The method of claim 25, further comprising determining the cellular or
35 biochemical function or both of said inhibitor protein.

50. The method of claim 25, wherein said identifying the bacterial target comprises identifying a phage-specific site of action.

5 51. An isolated, purified, or enriched nucleic acid sequence at least 15 nucleotides in length, wherein said sequence corresponds to at least a portion of a bacteriophage sequence, and wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

10

52. The nucleic acid sequence of claim 51, wherein said sequence comprises at least 50 nucleotides.

15 53. The nucleic acid sequence of claim 51, wherein said nucleic acid sequence corresponds to at least a portion of a nucleic acid sequence which encodes a product which provides a bacteria-inhibiting function.

54. The nucleic acid sequence of claim 53, wherein said nucleic acid sequence encodes a polypeptide which provides a bacteria-inhibiting function.

20

55. The nucleic acid sequence of claim 54, wherein said nucleic acid sequence is transcriptionally linked with regulatory sequences enabling induction of expression of said sequence.

25

56. An isolated, purified, or enriched polypeptide comprising at least a portion of a protein providing a bacteria-inhibiting function, wherein said polypeptide is normally encoded by a bacteriophage selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

30

57. The polypeptide of claim 56, wherein said polypeptide provides said bacteria-inhibiting function.

35 58. The polypeptide of claim 56, wherein said polypeptide comprises a portion at least 10 amino acid residues in length of a said polypeptide normally encoded by said bacteriophage.

59. A recombinant vector comprising a bacteriophage ORF corresponding to an ORF from a bacteriophage having a pathogenic bacterial host, wherein said
5 bacterial host is selected from the group consisting of uncharacterized bacteria of Table 1.

60. The vector of claim 59, wherein said vector is an expression vector.

10 61. The vector of claim 59, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage of Table 1.

62. The vector of claim 61, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD,
15 *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.

63. The vector of claim 60, wherein expression of said ORF is inducible.

20 64. A recombinant cell comprising a vector, wherein said vector comprises an ORF from a bacteriophage having a pathogenic bacterial host, wherein said bacterial host is selected from the group consisting of bacterial species of Table 1.

65. The recombinant cell of claim 64, wherein said bacteriophage is
25 selected from the group consisting of uncharacterized phage of Table 1.

66. The cell of claim 65, wherein said bacteriophage is selected from the group consisting of *Staphylococcus aureus* bacteriophage 77, 3A, 96, and 44AHJD, *Enterococcus* bacteriophage 182, and *Streptococcus pneumoniae* bacteriophage Dp-1.
30

67. The cell of claim 64, wherein said vector is an expression vector and expression of said ORF is inducible.

35 68. A method for identifying an antibacterial agent, comprising identifying an active portion of a product of a bacteria-inhibiting ORF of a bacteriophage.

69. The method of claim 68, further comprising constructing a synthetic peptidomimetic molecule, wherein the structure of said molecule corresponds to the structure of said active portion.

5

70. A method for identifying a compound active on a target of a bacteriophage inhibitor protein, comprising the step of contacting a bacterial target protein with a test compound; and determining whether said compound binds to or reduces the level of activity of said target protein,

10 wherein binding of said compound with said target protein or a reduction of the level of activity of said protein is indicative that said compound is active on said target and wherein said target is uncharacterized.

15 71. The method of claim 70, wherein said contacting is carried out *in vitro*.

72. The method of claim 70, wherein said contacting is carried out *in vivo* in a cell.

20 73. The method of claim 70, wherein said compound is a small molecule.

74. The method of claim 70, wherein said compound is a peptidomimetic compound.

25 75. The method of claim 70, wherein said compound is a fragment of a bacteriophage inhibitor protein.

76. The method of claim 70, further comprising determining the site of action of said compound on said target protein.

30

77. The method of claim 70, wherein said contacting is performed for a plurality of said target proteins.

35

78. A method of screening for potential antibacterial agents, comprising the step of determining whether any of a plurality of compounds is active on a target of a bacteriophage inhibitor protein,

wherein said target is naturally produced by a pathogenic bacterium.

79. The method of claim 78, wherein said plurality of compounds are small molecules.

5

80. The method of claim 78, wherein said determining is performed for a plurality of said targets.

10

81. A method for inhibiting a bacterium, comprising the step of; contacting said bacterium with a compound active on a target of a bacteriophage inhibitor protein, wherein said target or the target site is uncharacterized.

15

82. The method of claim 81, wherein said compound is said protein or an active fragment thereof.

83. The method of claim 81, wherein said compound is a structural mimetic of said protein.

20

84. The method of claim 81, wherein said compound is a small molecule.

85. The method of claim 81, wherein said contacting is performed *in vitro*.

25

86. The method of claim 81, wherein said contacting is performed *in vivo* in an animal.

87. The method of claim 86, wherein said animal is a human.

30

88. The method of claim 81, wherein said contacting is carried out *in vivo* in a plant.

89. The method of claim 81, wherein said bacterium is selected from the group of bacteria listed in Table 1.

35

90. A method for treating a bacterial infection in an animal suffering from an infection, comprising administering to said animal a therapeutically effective amount of compound active on a target of a bacteriophage inhibitor protein in a bacterium involved in said infection,

5 wherein said target is an uncharacterized target or the compound is active at an uncharacterized target site.

91. The method of claim 90, wherein said compound is a small molecule.

10 92. The method of claim 90, wherein said compound is a peptidomimetic compound.

93. The method of claim 90, wherein said compound is a fragment of a bacteriophage inhibitor protein.

15

94. The method of claim 90, wherein said animal is a mammal.

95. The method of claim 94, wherein said mammal is a human.

20 96. The method of claim 90, wherein said bacterium is selected from the group listed in Table 1.

97. The method of claim 90, wherein said bacteriophage inhibitor protein is from a bacteriophage selected from the group of bacteriophage listed in Table 1.

25

98. A method for prophylactically treating an animal at risk of an infection, comprising administering to said animal a prophylactically effective amount of a compound active on a target of a bacteriophage inhibitor protein,

30 wherein said target is an uncharacterized target or the site of action of said compound is an uncharacterized target site.

99. The method of claim 98, wherein said compound is a small molecule.

35 100. The method of claim 98, wherein said compound is a peptidomimetic compound.

101. The method of claim 98, wherein said compound is a fragment of a bacteriophage inhibitor protein.

102. The method of claim 98, wherein said animal is a mammal.

103. The method of claim 102, wherein said mammal is a human.

104. An antibacterial agent active on a target of a bacteriophage inhibitor protein, wherein said target is an uncharacterized target or said agent is active at a phage-specific site on said target.

105. The agent of claim 104, wherein said agent is a peptidomimetic of a bacteriophage inhibitor polypeptide.

106. The agent of claim 104, wherein said agent is a small molecule.

107. The agent of claim 104, wherein said agent is a fragment of a bacteriophage inhibitor polypeptide.

108. The agent of claim 104, wherein said agent is active at a phage-specific site on said target.

109. A method of making an antibacterial agent, comprising the steps of:

- a) identifying a target of a bacteriophage inhibitor polypeptide;
- b) screening a plurality of test compounds to identify a compound active on said target; and
- c) synthesizing said compound in an amount sufficient to provide a therapeutic effect when administered to an organism infected by a bacterium naturally producing said target.

110. The method of claim 109, wherein said compound is a small molecule.

111. The method of claim 109, wherein said compound is a peptidomimetic compound.

112. The method of claim 109, wherein said compound is a fragment or derivative of a bacteriophage inhibitor protein.

5 113. A computer readable device having recorded therein a nucleotide sequence of a portion of at least one bacteriophage genome of *Staphylococcus aureus* bacteriophage 77, bacteriophage 3A, or bacteriophage 96, a nucleotide sequence at least 95% identical to a said nucleotide sequence, a ribonucleic acid equivalent, a degenerate equivalent, a homologous sequence, or at least one amino acid sequence
10 encoded by said nucleotide sequence; and
 a nucleotide sequence or amino acid sequence analysis program,
 wherein said program can perform at least one sequence analysis on said nucleotide or amino acid sequence.

15 114. The device of claim 113, wherein said at least a portion of at least one bacteriophage genome comprises at least one ORF.

 115. The device of claim 113, wherein said device comprises a medium selected from the group consisting of floppy disk, computer hard drive, optical disk,
20 computer random access memory, and magnetic tape wherein said nucleotide or amino acid sequence or said program or both are recorded on said medium.

 116. The device of claim 113, wherein said portion of at least one bacteriophage genomic nucleotide sequence comprises at least 50% of at least one
25 bacteriophage genomic sequence.

 117. The device of claim 113, wherein said at least one bacteriophage nucleotide genomic sequence comprises portions of a plurality of bacteriophage nucleotide genomic sequences.

30

 118. A computer-based system for identifying biologically important portions of a bacteriophage genome, comprising:

 a) a data storage medium having recorded thereon a nucleotide sequence
35 corresponding to a portion of at least one bacteriophage genome, wherein said bacteriophage genome is uncharacterized;

- b) a set of instructions allowing searching of said sequence to analyze said sequence; and
- c) an output device.

5 119. The system of claim 118, wherein said output device comprises comprises a device selected from the group consisting of a printer, a video display, and a recording medium.

10 120. The system of claim 118, wherein said bacteriophage genome is of a bacteriophage selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

15 121. The system of claim 118, wherein said uncharacterized bacteriophage is selected from the group consisting of bacteriophage 77, 3A, and 96.

 122. A method for identifying or characterizing a bacteriophage ORF, comprising the steps of:

- a) providing a computer-based system for analyzing nucleic acid or amino acid sequence data, wherein said system comprises a data storage medium having recorded thereon at least one nucleotide or amino acid sequence corresponding to a portion of at least one uncharacterized bacteriophage genome, a set of instructions allowing searching of said sequence to analyze said sequence; and an output device;
- 20 b) analyzing at least a portion of at least one said sequence; and
- 25 c) outputting results of said analyzing to said output device.

 123. The method of claim 122, wherein said analysis identifies sequence similarity or homology with sequences selected from the group consisting of bacterial ORFs encoding products with related biological function; ORFs encoding known inhibitors or bacteria, essential bacterial ORFs.

30

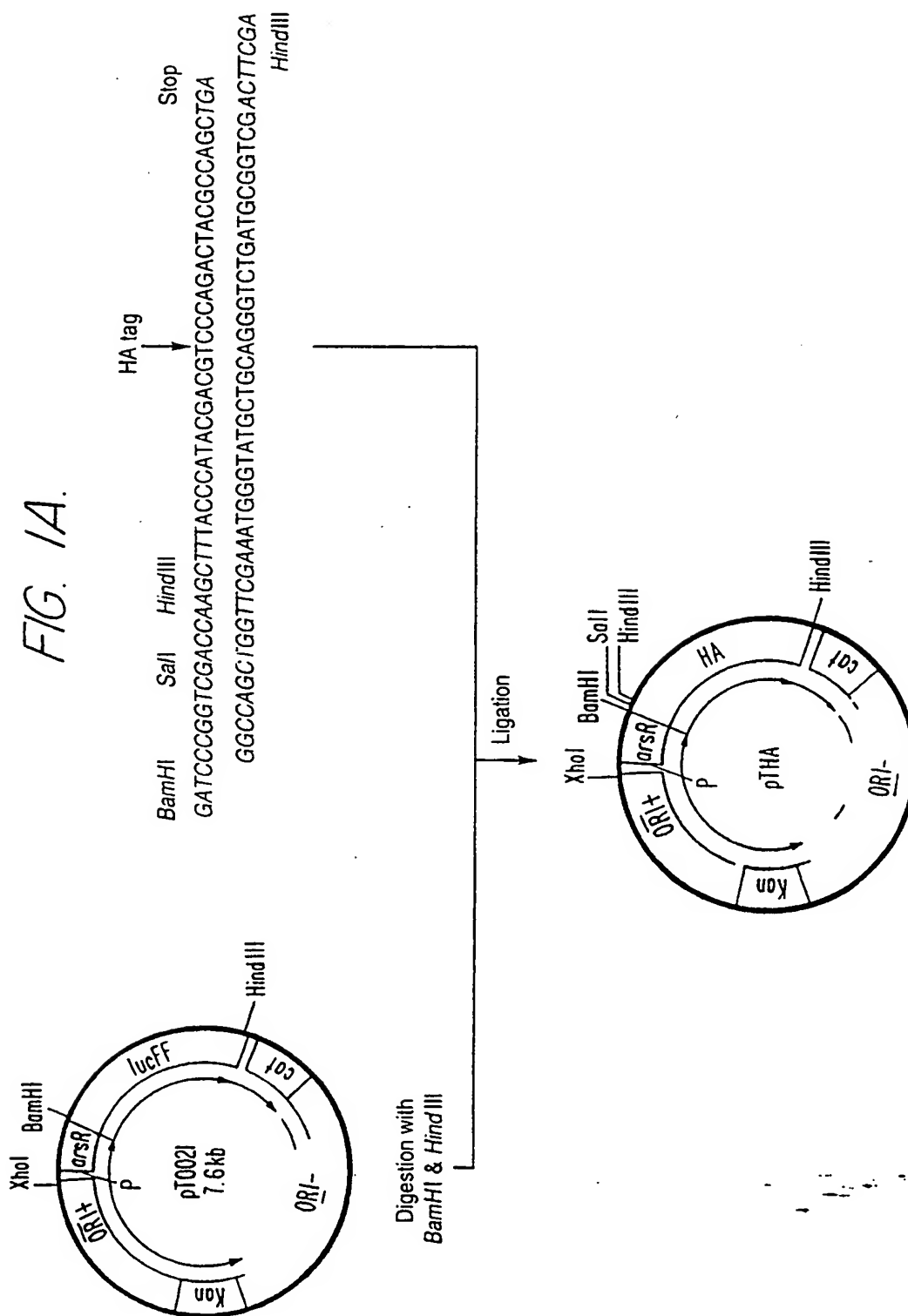
 124. The method of claim 122, wherein said analysis comprises identifying a probable biological function based on identification of structural elements or sequence homology or similarity.

35

 125. The method of claim 122, wherein said bacteriophage is selected from the group consisting of uncharacterized bacteriophage listed in Table 1.

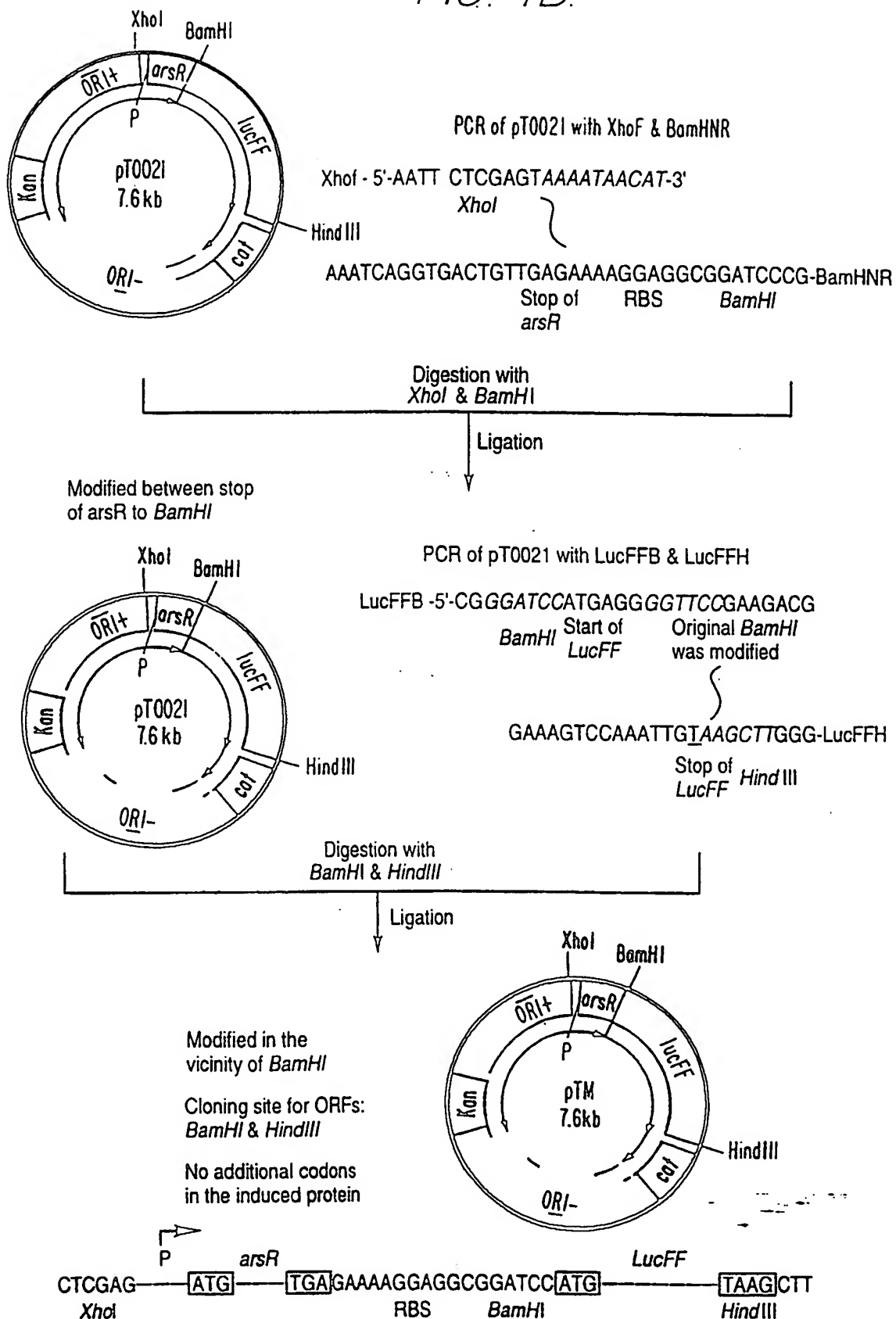
126. The method of claim 125, wherein said uncharacterized bacteriophage is selected from bacteriophage 77, 3A, and 96.

01/11



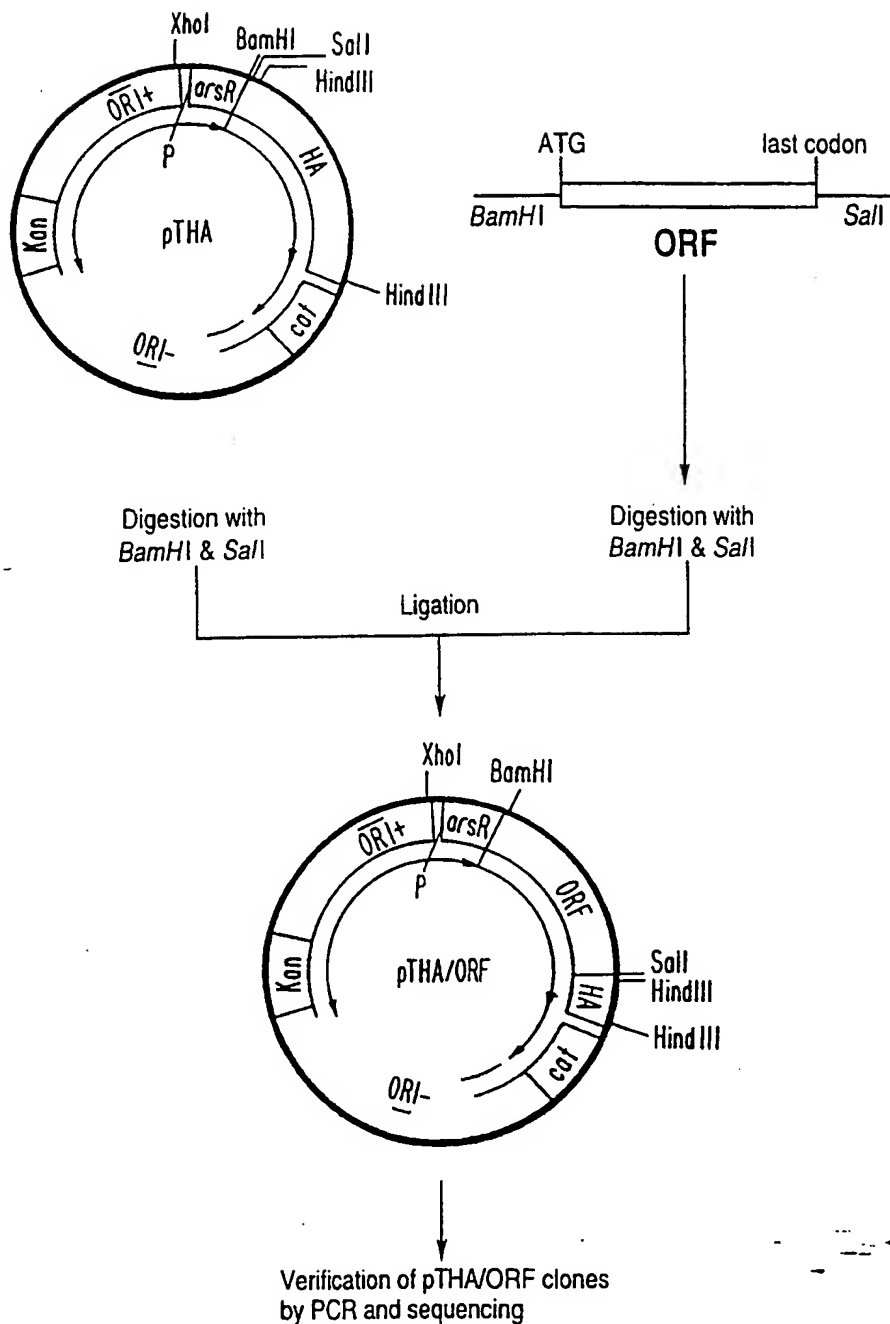
02/11

FIG. 1B.



03/11

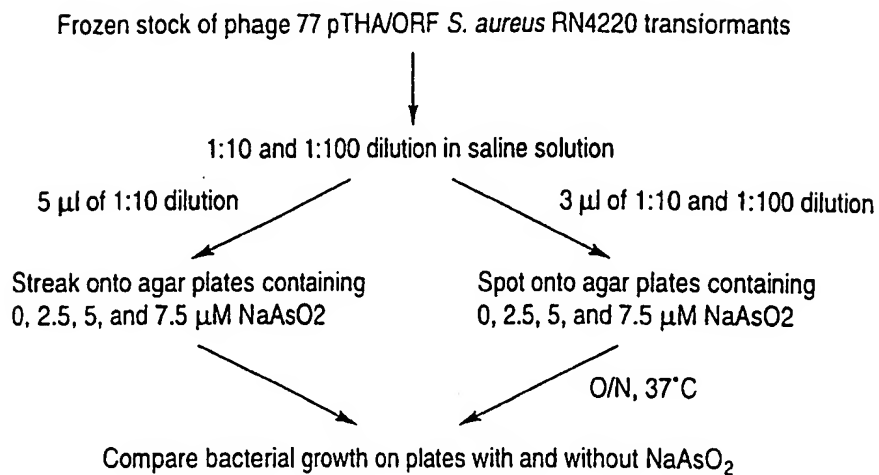
FIG. 2.



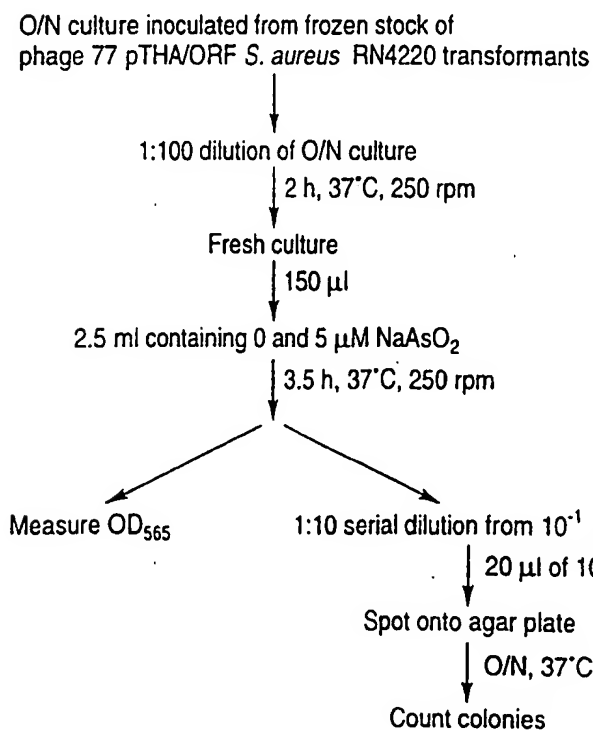
04/11

FIG. 3.

(A) Functional assay on semi-solid support media



(B) Functional assay in liquid medium

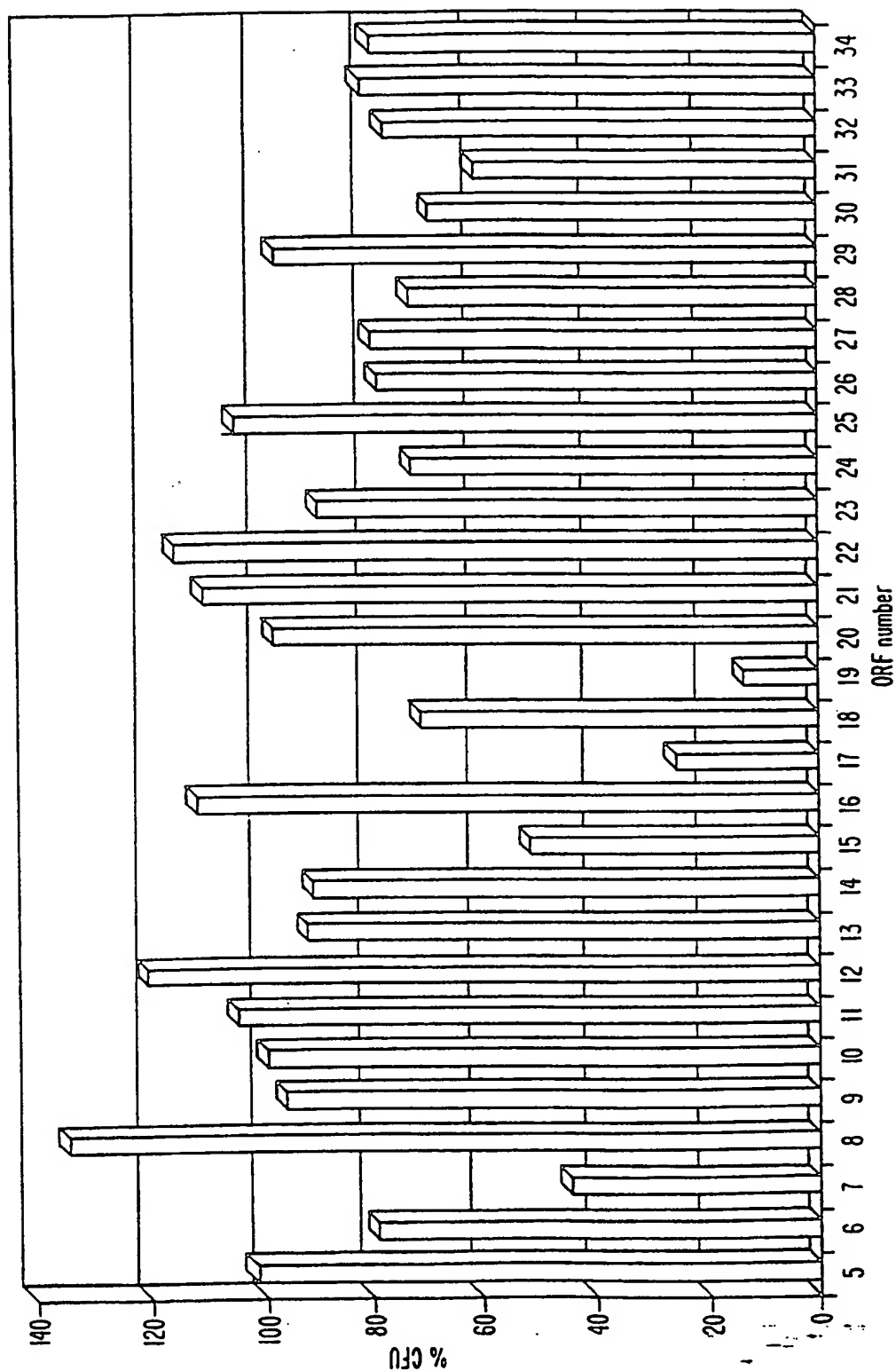


SUBSTITUTE SHEET (RULE 26)

05/11

A. Inhibition of bacterial growth with individual ORFs of a S. aureus Bacteriophage.

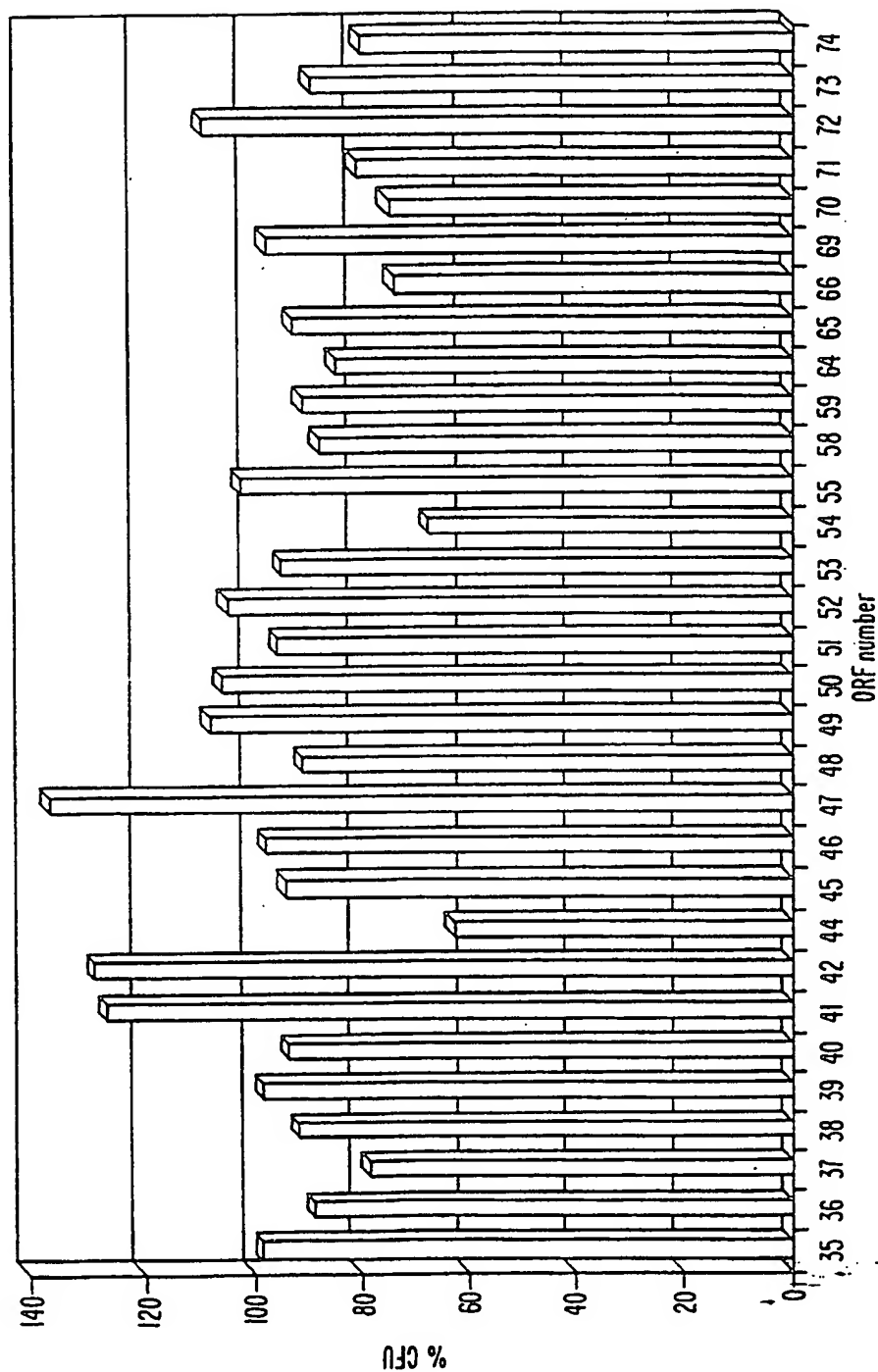
FIG. 4A.



06/11

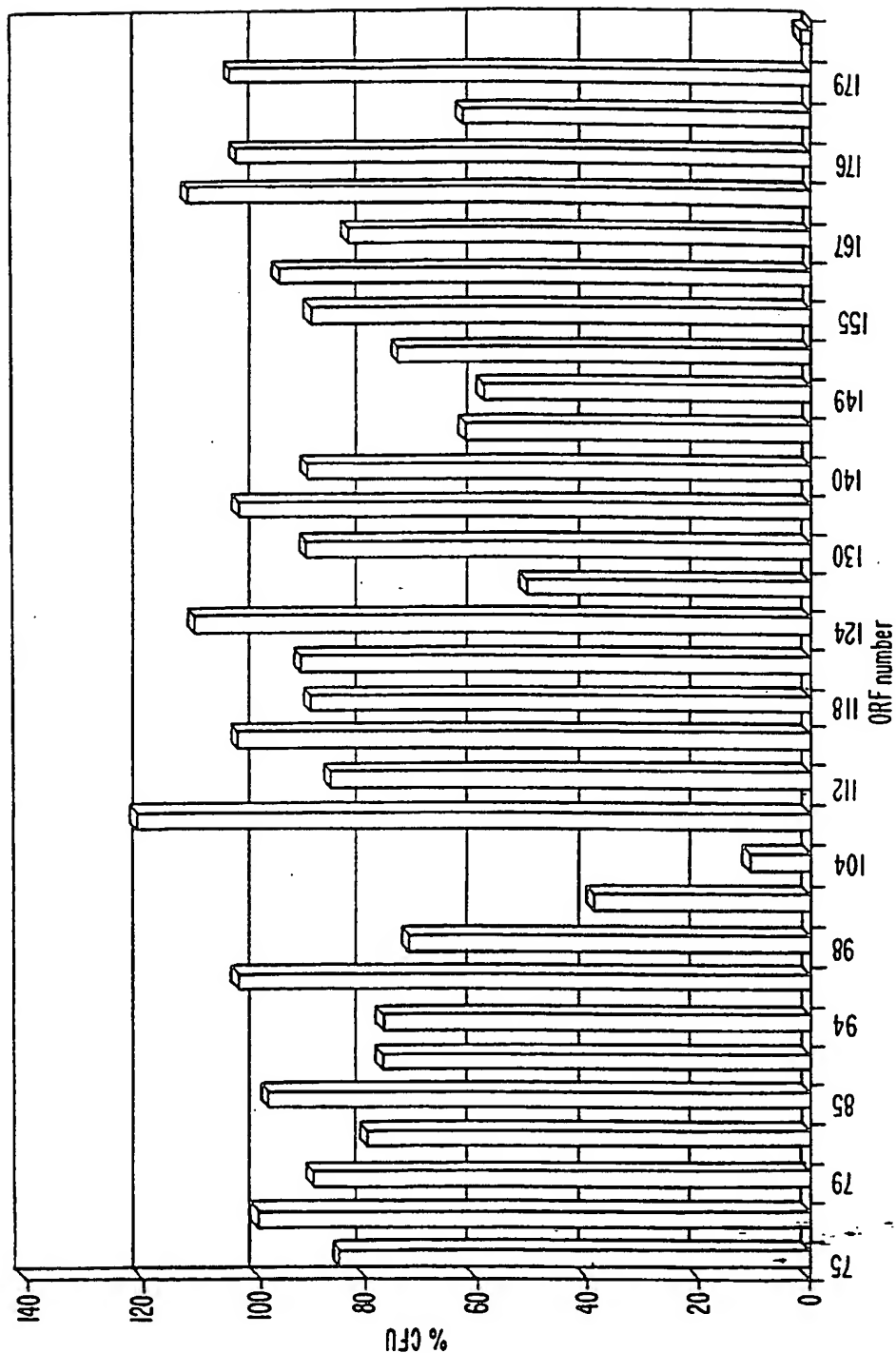
B. Inhibition of bacterial growth with individual ORFs of a *S. aureus* Bacteriophage.

FIG. 4B.



07/11

FIG. 4C. C. Inhibition of bacterial growth with individual ORFs of a *S. aureus* Bacteriophage.



08/11

FIG. 5.

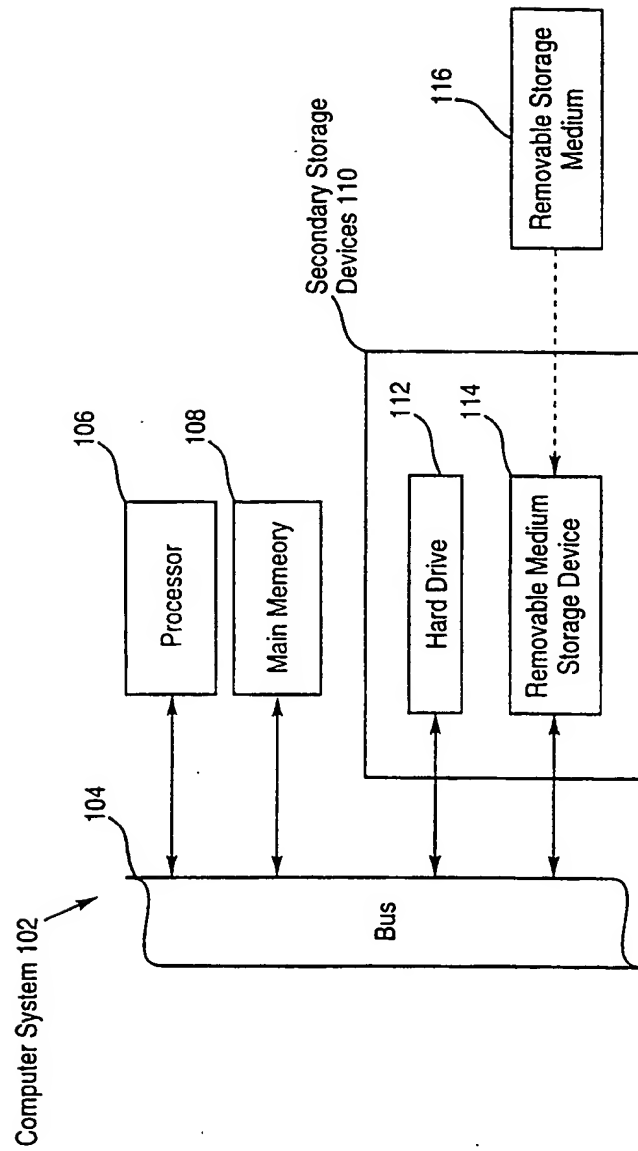
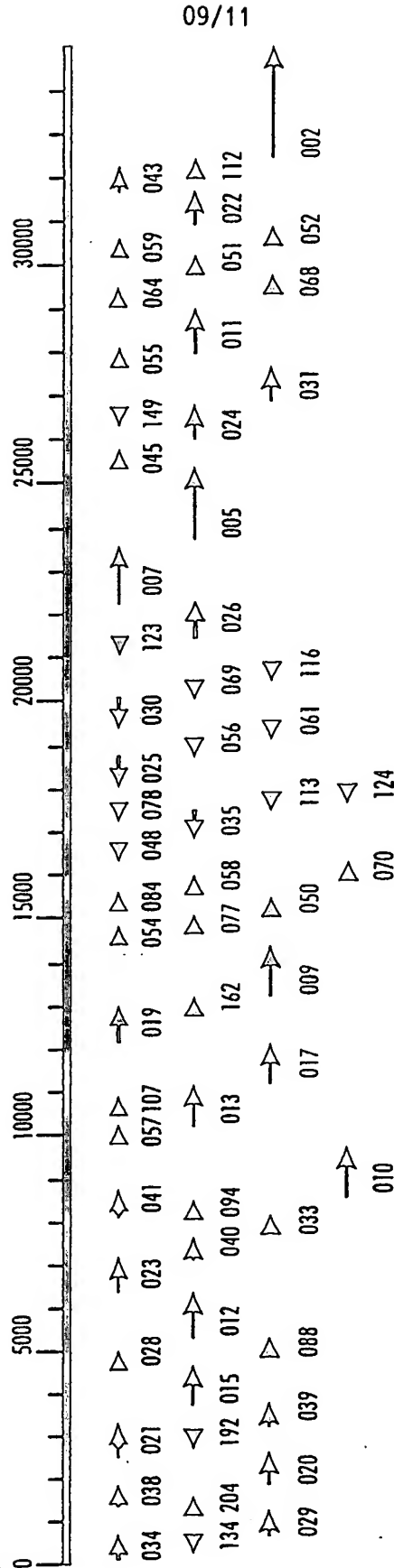


Fig 6A Fig 6B

Fig. 6A

PHAGE: BACTERIOPHAGE Dp1
MINIMAL ORF SIZE: 33 A. A.
ORFS "WITH" RBS.
NUMBER OF ORFS: 85

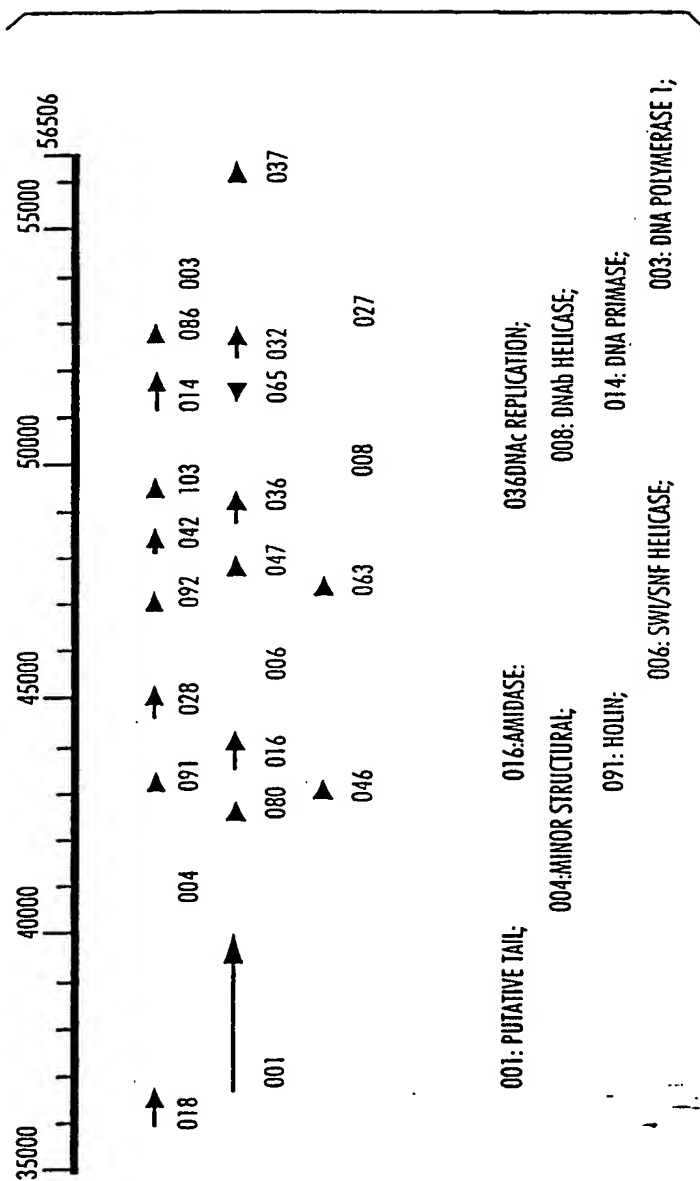


029: exsB; 012: DNA pol. III beta;
038: exsC; 6-PYRUVYL-TETRAHYDROPTERIN
020: exsD; COENZYME PQQ; 013: DNA POL III GAMMA AND TAU;
021: GTP CYCLOHYDROLASE;
039: CITRULLINE BIOSYNTHESIS;
041: dUTPASE;
010: RecA;

007: TERMINASE; 011: MAJOR HEAD; 002: TAIL;

10/11

Fig. 6B



11/11

FIG. 7.

Abbreviations:

kan: gene encoding kanamycin resistance
 cat: gene encoding chloramphenicol resistance
 ori + and -: origin of replication in gram-positive and
 gram-negative bacteria, respectively
 arsR: gene encoding regulatory protein of the ars promoter
 P: ars promoter
 lucFF: gene encoding luciferase protein. This portion will
 be removed and replaced by individual *S. aureus* phage
 genes.

Reference:

Tauriainen et al., Appl. Environ. Microbio. 1997. 63: 4456-4461

